

97965

STIC-Biotech/ChemLib

From: Fredman, Jeffrey
Sent: Wednesday, July 02, 2003 12:51 PM
To: STIC-Biotech/ChemLib
Cc: Schultz, James
Subject: FW: Rush Sequence search request 09/780,929

PLEASE RUSH.

I Approve.

Jeff Fredman

CRF

-----Original Message-----

From: Schultz, James
Sent: Wednesday, July 02, 2003 9:38 AM
To: Fredman, Jeffrey
Subject: Rush Sequence search request 09/780,929

Dear Jeff,

Would you please approve the rush sequence search below? This case has already been searched and is ready for allowance, but my SPE wants it searched one more way before we pass it out.

Thanks,

Doug Schultz

Dear STIC-biotech searchers,

Could you please run a length limited nucleotide sequence search on SEQ ID NOS 97 (15 nt long) and 98 (18 nt long) in the above entitled case, where the maximum size of the returned hit is no longer than 60 nucleotides? I need both sequences searched in the **interference** databases as well.

Thanks,

Doug Schultz

J. Douglas Schultz, Ph.D.
AU 1635 (Biotechnology)
Patent Examiner
United States Patent and Trademark Office
(703) 308-9355
(703) 746-3973 (fax)
Office: CM1 12E18
Mail: CM1 11E12

Searcher: _____
Phone: _____
Location: _____
Date Picked Up: _____
Searcher Prep/Review: _____
Clerical: _____
Online time: _____

TYPE OF SEARCH:

NA Sequences: _____
AA Sequences: _____
Structures: _____
Bibliographic: _____
Litigation: _____
Full text: _____
Patent Family: _____
Other: _____

VENDOR/COST (where applic.)

STN: _____
DIALOG: _____
Questel/Orbit: _____
DRLink: _____
Lexis/Nexis: _____
Sequence Sys.: _____
WWW/Internet: _____
Other (specify): _____

GenCore version 5.1.6
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: July 6, 2003, 14:51:21 ; Search time 520.455 seconds

(without alignments)
206.249 Million cell updates/sec

Title: US-09-780-929-97

Perfect score: 15

Sequence: 1 agauaacgugaagau 15

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 8255821 seqs, 3578102051 residues

Total number of hits satisfying chosen parameters: 9359164

Minimum DB seq length: 0

Maximum DB seq length: 60

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

Pending_Patents_NA_New:*
1: /cgn2_6/ptodata/2/pna/PCT_NEW_COMB.seq:*
2: /cgn2_6/ptodata/2/pna/PCT_NEW_COMB.seq4:*
3: /cgn2_6/ptodata/2/pna/US06_NEW_COMB.seq:*
4: /cgn2_6/ptodata/2/pna/US06_NEW_COMB.seq4:*
5: /cgn2_6/ptodata/2/pna/US07_NEW_COMB.seq:*
6: /cgn2_6/ptodata/2/pna/US07_NEW_COMB.seq4:*
7: /cgn2_6/ptodata/2/pna/US08_NEW_COMB.seq:*
8: /cgn2_6/ptodata/2/pna/US08_NEW_COMB.seq4:*
9: /cgn2_6/ptodata/2/pna/US09_NEW_COMB.seq:*
10: /cgn2_6/ptodata/2/pna/US09_NEW_COMB.seq4:*
11: /cgn2_6/ptodata/2/pna/US09_NEW_COMB.seq3:*
12: /cgn2_6/ptodata/2/pna/US09_NEW_COMB.seq4:*
13: /cgn2_6/ptodata/2/pna/US10_NEW_COMB.seq:*
14: /cgn2_6/ptodata/2/pna/US10_NEW_COMB.seq4:*
15: /cgn2_6/ptodata/2/pna/US10_NEW_COMB.seq3:*
16: /cgn2_6/ptodata/2/pna/US10_NEW_COMB.seq4:*
17: /cgn2_6/ptodata/2/pna/US60_NEW_COMB.seq:*
18: /cgn2_6/ptodata/2/pna/US60_NEW_COMB.seq3:*
19: /cgn2_6/ptodata/2/pna/US60_NEW_COMB.seq4:*
20: /cgn2_6/ptodata/2/pna/US60_NEW_COMB.seq3:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
C 1	13.4	89.3	25	14	US-10-355-577-24855
C 2	13.4	89.3	25	18	US-60-427-836-243678
C 3	13	86.7	25	14	US-10-355-577-179244
C 4	12.4	82.7	25	18	US-60-427-836-118114
C 5	12.4	82.7	23	14	US-10-310-188-75880
C 6	12.4	82.7	25	10	US-09-560-222-72811
C 7	12.4	82.7	25	10	US-09-560-222-72812
C 8	12.4	82.7	25	11	US-09-953-570-48322
C 9	12.4	82.7	25	11	US-09-953-570-48329
C 10	12.4	82.7	25	11	US-09-954-445A-10200
C 11	12.4	82.7	25	11	US-09-954-445A-16612
C 12	12.4	82.7	25	11	US-09-954-445A-16620
C 13	12.4	82.7	25	11	US-09-954-445A-50048

14	12.4	82.7	25	14	US-10-355-577-40591	Sequence 40591, A
15	12.4	82.7	25	14	US-10-355-577-366476	Sequence 366476, A
16	12.4	82.7	25	14	US-10-355-577-485354	Sequence 485354, A
17	12.4	82.7	25	14	US-10-355-577-511942	Sequence 511942, A
18	12.4	82.7	25	14	US-10-355-577-735345	Sequence 735345, A
19	12.4	82.7	25	14	US-10-355-577-834196	Sequence 834196, A
20	12.4	82.7	25	14	US-10-355-577-907847	Sequence 907847, A
21	12.4	82.7	25	18	US-60-427-808-475949	Sequence 475949, A
22	12.4	82.7	25	18	US-60-427-808-568077	Sequence 568077, A
23	12.4	82.7	25	18	US-60-427-808-695630	Sequence 695630, A
24	12.4	82.7	25	18	US-60-427-808-928248	Sequence 928248, A
25	12.4	82.7	25	18	US-60-427-836-166264	Sequence 166264, A
26	12.4	82.7	25	18	US-60-427-836-525766	Sequence 525766, A
27	12.4	82.7	25	18	US-60-427-836-674310	Sequence 674310, A
28	12.4	82.7	25	19	US-60-469-545-95587	Sequence 95587, A
29	12.4	82.7	25	19	US-60-469-545-96568	Sequence 96568, A
30	12.4	82.7	25	19	US-60-469-545-141609	Sequence 141609, A
31	12.4	82.7	25	19	US-60-469-545-142590	Sequence 142590, A
32	12.4	82.7	25	20	US-60-475-871-165169	Sequence 165169, A
33	12.4	82.7	41	15	US-10-408-085-133	Sequence 133, App
34	12.4	82.7	41	15	US-10-408-085-359	Sequence 359, App
35	12	80.0	22	14	US-10-310-188-49187	Sequence 49187, A
36	12	80.0	25	10	US-09-660-222-100324	Sequence 100324, A
37	12	80.0	25	11	US-09-953-570-45866	Sequence 45866, A
38	12	80.0	25	11	US-09-953-570-45892	Sequence 45892, A
39	12	80.0	25	14	US-10-098-263B-27903	Sequence 27903, A
40	12	80.0	25	14	US-10-355-577-53971	Sequence 53971, A
41	12	80.0	25	14	US-10-355-577-143487	Sequence 143487, A
42	12	80.0	25	14	US-10-355-577-518580	Sequence 518580, A
43	12	80.0	25	14	US-10-355-577-653875	Sequence 653875, A
44	12	80.0	25	14	US-10-355-577-827041	Sequence 827041, A
45	12	80.0	25	14	US-10-355-577-899636	Sequence 899636, A

ALIGNMENTS

RESULT 1
US-10-355-577-24855/c
; Sequence 24855, Application US/10355577
; GENERAL INFORMATION:
; APPLICANT: Mitmann, Michael
; TITLE OF INVENTION: Methods of Genetic Analysis of Probes: HG-U133
; FILE REFERENCE: 3121
; CURRENT APPLICATION NUMBER: US/10/355,577
; CURRENT FILING DATE: 2003-01-31
; NUMBER OF SEQ ID NOS: 997516
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 24855
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapien
US-10-355-577-24855

Query Match 89.3%; Score 13.4; DB 14; Length 25;
Best Local Similarity 73.3%; Pred. No. 1.9e+03;
Matches 11; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

QY 1 AGAUACGUGAAGAU 15
|||:||||:| |||:
Db 25 AGATAAGTCGAGAT 11

RESULT 2
US-60-427-836-243678/c
; Sequence 243678, Application US/60427836
; GENERAL INFORMATION:
; APPLICANT: Xue Mei Zhou
; TITLE OF INVENTION: Methods of Genetic Analysis of Rat
; FILE REFERENCE: 3527
; CURRENT APPLICATION NUMBER: US/60/427,836
; CURRENT FILING DATE: 2002-11-20
; NUMBER OF SEQ ID NOS: 699466

; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 243678
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Rattus norvegicus
US-60-427-836-243678

Query Match 89.3%; Score 13.4; DB 18; Length 25;
Best Local Similarity 73.3%; Pred. No. 1.9e+03;
Matches 11; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

QY 1 AGAUAACGUGAAGAU 15
| | | | | | | | | | | | | | | | | | | | | |
Db 21 AGATAAAGTGAAGAT 7

RESULT 3
US-10-355-577-179244/c
; Sequence 179244, Application US/10355577
; GENERAL INFORMATION:
; APPLICANT: Mittmann, Michael
; TITLE OF INVENTION: Methods of Genetic Analysis of Probes: HG-U133
; FILE REFERENCE: 3121
; CURRENT APPLICATION NUMBER: US/10/355,577
; CURRENT FILING DATE: 2003-01-31
; NUMBER OF SEQ ID NOS: 997516
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 179244
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapien
US-10-355-577-179244

Query Match 86.7%; Score 13; DB 14; Length 25;
Best Local Similarity 84.6%; Pred. No. 3.1e+03;
Matches 11; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 2 GAUAACGUGAAGA 14
| | | | | | | | | | | | | | | | | | | | | |
Db 22 GATAACGTGAAGA 10

RESULT 4
US-60-427-836-118114
; Sequence 118114, Application US/60427836
; GENERAL INFORMATION:
; APPLICANT: Xue Mei Zhou
; TITLE OF INVENTION: Methods of Genetic Analysis of Rat
; FILE REFERENCE: 3527
; CURRENT APPLICATION NUMBER: US/60/427,836
; CURRENT FILING DATE: 2002-11-20
; NUMBER OF SEQ ID NOS: 699466
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 118114
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Rattus norvegicus
US-60-427-836-118114

Query Match 86.7%; Score 13; DB 18; Length 25;
Best Local Similarity 84.6%; Pred. No. 3.1e+03;
Matches 11; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGAUAACGUGAAG 13
| | | | | | | | | | | | | | | | | | | | | |
Db 1 AGATAACGTGAAG 13

RESULT 5
US-10-310-188-75880/c
; Sequence 75880, Application US/10310188
; GENERAL INFORMATION:
; APPLICANT: RosettaGenomics

; TITLE OF INVENTION: BIOINFORMATICAALLY DETECTABLE GROUP OF NOVEL VIRAL REGULATORY
; TITLE OF INVENTION: USES THEREOF
; FILE REFERENCE: 47487
; CURRENT APPLICATION NUMBER: US/10/310,188
; CURRENT FILING DATE: 2002-12-19
; NUMBER OF SEQ ID NOS: 86841
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 75880
; LENGTH: 23
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-310-188-75880

Query Match 82.7%; Score 12.4; DB 14; Length 23;
Best Local Similarity 78.6%; Pred. No. 6.7e+03;
Matches 11; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 1 AGAUAACGUGAAGA 14
| | | | | | | | | | | | | | | | | | | | | |
Db 23 AAATAACGTGAAGA 10

RESULT 6
US-09-660-222-72811/c
; Sequence 72811, Application US/09660222
; GENERAL INFORMATION:
; APPLICANT: Mittmann et al.
; TITLE OF INVENTION: Methods of Genetic Analysis of Human
; FILE REFERENCE: 3102.1
; CURRENT APPLICATION NUMBER: US/09/660,222
; CURRENT FILING DATE: 2000-09-12
; PRIOR APPLICATION NUMBER: 60/164,973
; PRIOR FILING DATE: 1999-11-11
; NUMBER OF SEQ ID NOS: 140981
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 72811
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo Sapiens
; PUBLICATION INFORMATION:
; DATABASE ACCESSION NUMBER: GenBank S80343
US-09-660-222-72811

Query Match 82.7%; Score 12.4; DB 10; Length 25;
Best Local Similarity 78.6%; Pred. No. 6.7e+03;
Matches 11; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 1 AGAUAACGUGAAGA 14
| | | | | | | | | | | | | | | | | | | | | |
Db 16 AGATAACGTGCAGA 3

RESULT 7
US-09-660-222-72812/c
; Sequence 72812, Application US/09660222
; GENERAL INFORMATION:
; APPLICANT: Mittmann et al.
; APPLICANT: Affymetrix, Inc.
; TITLE OF INVENTION: Methods of Genetic Analysis of Human
; FILE REFERENCE: 3102.1
; CURRENT APPLICATION NUMBER: US/09/660,222
; CURRENT FILING DATE: 2000-09-12
; PRIOR APPLICATION NUMBER: 60/164,973
; PRIOR FILING DATE: 1999-11-11
; NUMBER OF SEQ ID NOS: 140981
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 72812
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo Sapiens
; PUBLICATION INFORMATION:
; DATABASE ACCESSION NUMBER: GenBank S80343

US-09-660-222-72812

Query Match 82.7%; Score 12.4; DB 10; Length 25;
Best Local Similarity 78.6%; Pred. No. 6.7e+03;
Matches 11; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 1 AGAUAACGUGAAGA 14
|||:||||:|
DB 22 AGATAACGTGCAGA 9

RESULT 8

US-09-570-48322/c
; Sequence 48322, Application US/09953570
; GENERAL INFORMATION:
; APPLICANT: Mittmann, Michael
; TITLE OF INVENTION: Methods of Genetic Analysis of Yeast
; FILE REFERENCE: 3110.1
; CURRENT APPLICATION NUMBER: US/09/953,570
; CURRENT FILING DATE: 2001-09-13
; PRIOR APPLICATION NUMBER: 60/232,638
; PRIOR FILING DATE: 2000-09-14
; NUMBER OF SEQ ID NOS: 138410
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 48322
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Saccharomyces Cerevisiae
US-09-570-48322

Query Match 82.7%; Score 12.4; DB 11; Length 25;
Best Local Similarity 78.6%; Pred. No. 6.7e+03;
Matches 11; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 1 AGAUAACGUGAAGA 14
|||:||||:|
DB 14 AGATAACGTGTAGA 1

RESULT 9

US-09-570-48329/c
; Sequence 48329, Application US/09953570
; GENERAL INFORMATION:
; APPLICANT: Mittmann, Michael
; TITLE OF INVENTION: Methods of Genetic Analysis of Yeast
; FILE REFERENCE: 3110.1
; CURRENT APPLICATION NUMBER: US/09/953,570
; CURRENT FILING DATE: 2001-09-13
; PRIOR APPLICATION NUMBER: 60/232,638
; PRIOR FILING DATE: 2000-09-14
; NUMBER OF SEQ ID NOS: 138410
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 48329
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Saccharomyces Cerevisiae
US-09-570-48329

Query Match 82.7%; Score 12.4; DB 11; Length 25;
Best Local Similarity 78.6%; Pred. No. 6.7e+03;
Matches 11; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 1 AGAUAACGUGAAGA 14
|||:||||:|
DB 20 AGATAACGTGTAGA 7

RESULT 10

US-09-954-445A-10200/c
; Sequence 10200, Application US/09954445A
; GENERAL INFORMATION:
; APPLICANT: Mittmann, Michael
; TITLE OF INVENTION: Methods of Genetic Analysis of Arabidopsis Thaliana

; FILE REFERENCE: 3116.1
; CURRENT APPLICATION NUMBER: US/09/954,445A
; CURRENT FILING DATE: 2000-09-17
; PRIOR APPLICATION NUMBER: 60/233,620
; PRIOR FILING DATE: 2000-09-18
; NUMBER OF SEQ ID NOS: 131820
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 10200
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Arabidopsis thaliana
US-09-954-445A-10200

Query Match 82.7%; Score 12.4; DB 11; Length 25;
Best Local Similarity 78.6%; Pred. No. 6.7e+03;
Matches 11; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 1 AGAUAACGUGAAGA 14
|||:||||:|
DB 17 AGATAACGTGAAGA 4

RESULT 11

US-09-954-445A-16612
; Sequence 16612, Application US/09954445A
; GENERAL INFORMATION:
; APPLICANT: Mittmann, Michael
; TITLE OF INVENTION: Methods of Genetic Analysis of Arabidopsis Thaliana
; FILE REFERENCE: 3116.1
; CURRENT APPLICATION NUMBER: US/09/954,445A
; CURRENT FILING DATE: 2000-09-17
; PRIOR APPLICATION NUMBER: 60/233,620
; PRIOR FILING DATE: 2000-09-18
; NUMBER OF SEQ ID NOS: 131820
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 16612
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Arabidopsis thaliana
US-09-954-445A-16612

Query Match 82.7%; Score 12.4; DB 11; Length 25;
Best Local Similarity 71.4%; Pred. No. 6.7e+03;
Matches 10; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

QY 2 GAUAACGUGAAGAU 15
||:||||:|
DB 6 GATAACGTGAATAT 19

RESULT 12

US-09-954-445A-16620
; Sequence 16620, Application US/09954445A
; GENERAL INFORMATION:
; APPLICANT: Mittmann, Michael
; TITLE OF INVENTION: Methods of Genetic Analysis of Arabidopsis Thaliana
; FILE REFERENCE: 3116.1
; CURRENT APPLICATION NUMBER: US/09/954,445A
; CURRENT FILING DATE: 2000-09-17
; PRIOR APPLICATION NUMBER: 60/233,620
; PRIOR FILING DATE: 2000-09-18
; NUMBER OF SEQ ID NOS: 131820
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 16620
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Arabidopsis thaliana
US-09-954-445A-16620

Query Match 82.7%; Score 12.4; DB 11; Length 25;
Best Local Similarity 71.4%; Pred. No. 6.7e+03;
Matches 10; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

QY 2 GAUAACGUGAAGAU 15
||:||||:|||||:
Db 1 GATAACGTGATAT 14

RESULT 13

US-09-954-445A-50048
; Sequence 50048, Application US/09954445A
; GENERAL INFORMATION:
; APPLICANT: Mittmann, Michael
; TITLE OF INVENTION: Methods of Genetic Analysis of Arabidopsis Thaliana
; FILE REFERENCE: 3116.1
; CURRENT APPLICATION NUMBER: US/09/954.445A
; CURRENT FILING DATE: 2000-09-17
; PRIOR APPLICATION NUMBER: 60/233,620
; PRIOR FILING DATE: 2000-09-18
; NUMBER OF SEQ ID NOS: 131820
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 50048
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Arabidopsis thaliana
US-09-954-445A-50048

Query Match 82.7%; Score 12.4; DB 11; Length 25;
Best Local Similarity 71.4%; Pred. No. 6.7e+03;
Matches 10; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

QY 2 GAUAACGUGAAGAU 15
||:||||:|||||:
Db 6 GATTACGTGATAT 19

RESULT 14

US-10-355-577-40591
; Sequence 40591, Application US/10355577
; GENERAL INFORMATION:
; APPLICANT: Mittmann, Michael
; TITLE OF INVENTION: Methods of Genetic Analysis of Probes: HG-U133
; FILE REFERENCE: 3121
; CURRENT APPLICATION NUMBER: US/10/355.577
; CURRENT FILING DATE: 2003-01-31
; NUMBER OF SEQ ID NOS: 997516
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 40591
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapien
US-10-355-577-40591

Query Match 82.7%; Score 12.4; DB 14; Length 25;
Best Local Similarity 85.7%; Pred. No. 6.7e+03;
Matches 12; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 AGAUAACGUGAAGA 14
||| ||||:|||||
Db 2 AGAGAACGTGAAGA 15

RESULT 15

US-10-355-577-366476/c
; Sequence 366476, Application US/10355577
; GENERAL INFORMATION:
; APPLICANT: Mittmann, Michael
; TITLE OF INVENTION: Methods of Genetic Analysis of Probes: HG-U133
; FILE REFERENCE: 3121
; CURRENT APPLICATION NUMBER: US/10/355.577
; CURRENT FILING DATE: 2003-01-31
; NUMBER OF SEQ ID NOS: 997516
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 366476
; LENGTH: 25
; TYPE: DNA

; ORGANISM: Homo sapien
US-10-355-577-366476

Query Match 82.7%; Score 12.4; DB 14; Length 25;
Best Local Similarity 78.6%; Pred. No. 6.7e+03;
Matches 11; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 1 AGAUAACGUGAAGA 14
|||:||||:|||||
Db 18 AGATAACGTGCAGA 5

Search completed: July 6, 2003, 16:49:13
Job time : 522.455 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2003 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: July 6, 2003, 14:26:16 ; Search time 493.182 seconds
(without alignments)
885.154 Million cell updates/sec

Title: US-09-780-929-97
Perfect score: 15
Sequence: 1 agauaacgugaagau 15

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 2054640 seqs, 14551402878 residues
Total number of hits satisfying chosen parameters: 897812

Minimum DB seq length: 0
Maximum DB seq length: 60

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :

GenEmbl.*

- 1: gb_ba.*
- 2: gb_htg.*
- 3: gb_in.*
- 4: gb_om.*
- 5: gb_ov.*
- 6: gb_pat.*
- 7: gb_ph.*
- 8: gb_pl.*
- 9: gb_pr.*
- 10: gb_ro.*
- 11: gb_sts.*
- 12: gb_sy.*
- 13: gb_un.*
- 14: gb_vi.*
- 15: em_ba.*
- 16: em_fun.*
- 17: em_hum.*
- 18: em_in.*
- 19: em_mu.*
- 20: em_om.*
- 21: em_or.*
- 22: em_ov.*
- 23: em_pat.*
- 24: em_ph.*
- 25: em_pl.*
- 26: em_ro.*
- 27: em_sts.*
- 28: em_un.*
- 29: em_vi.*
- 30: em_htg_hum.*
- 31: em_htg_inv.*
- 32: em_htg_other.*
- 33: em_htg_mus.*
- 34: em_htg_pln.*
- 35: em_htg_rod.*
- 36: em_htg_mam.*
- 37: em_htg_vrt.*
- 38: em_sy.*
- 39: em_htgo_hum.*
- 40: em_htgo_mus.*
- 41: em_htgo_other.*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	DB	ID	Description
1	15	100.0	15	6	AX214295	Sequence
2	15	100.0	27	6	AX214239	Sequence
3	15	100.0	28	6	AX214237	Sequence
4	15	100.0	28	6	AX214238	Sequence
5	15	100.0	28	6	AX214240	Sequence
6	15	100.0	28	6	AX214241	Sequence
7	15	100.0	28	6	AX214242	Sequence
8	15	100.0	28	6	AX214243	Sequence
9	15	100.0	28	6	AX214244	Sequence
10	15	100.0	28	6	AX214245	Sequence
11	15	100.0	28	6	AX214246	Sequence
12	15	100.0	28	6	AX214247	Sequence
13	15	100.0	28	6	AX214248	Sequence
14	15	100.0	28	6	AX214249	Sequence
15	15	100.0	28	6	AX214250	Sequence
16	15	100.0	28	6	AX214251	Sequence
17	15	100.0	28	6	AX214252	Sequence
18	15	100.0	28	6	AX214253	Sequence
19	15	100.0	28	6	AX214254	Sequence
20	15	100.0	28	6	AX214255	Sequence
21	15	100.0	28	6	AX214256	Sequence
22	15	100.0	28	6	AX214257	Sequence
23	15	100.0	28	6	AX214258	Sequence
24	15	100.0	28	6	AX214259	Sequence
25	15	100.0	28	6	AX214260	Sequence
26	15	100.0	28	6	AX214261	Sequence
27	15	100.0	28	6	AX214262	Sequence
28	15	100.0	28	6	AX214263	Sequence
29	15	100.0	28	6	AX214264	Sequence
30	15	100.0	28	6	AX214265	Sequence
31	15	100.0	28	6	AX214266	Sequence
32	15	100.0	28	6	AX214267	Sequence
33	15	100.0	28	6	AX214268	Sequence
34	15	100.0	28	6	AX214269	Sequence
35	15	100.0	28	6	AX214270	Sequence
36	15	100.0	28	6	AX214271	Sequence
37	15	100.0	28	6	AX214272	Sequence
38	15	100.0	28	6	AX214273	Sequence
39	15	100.0	28	6	AX214274	Sequence
40	15	100.0	28	6	AX214275	Sequence
41	15	100.0	28	6	AX214276	Sequence
42	15	100.0	28	6	AX214277	Sequence
43	15	100.0	28	6	AX214278	Sequence
44	15	100.0	28	6	AX214279	Sequence
45	15	100.0	28	6	AX214280	Sequence

ALIGNMENTS

RESULT 1	AX214295	AX214295	15 bp	mrna	linear	PAT 06-SEP-2001
LOCUS	Sequence	108 from Patent WO0159102.				
DEFINITION	AX214295					
ACCESSION	AX214295					
VERSION	AX214295.1	GI:15524372				
KEYWORDS						
SOURCE		synthetic construct.				
ORGANISM		synthetic construct				
REFERENCE		1 (bases 1 to 15)				
AUTHORS		Breaker, R. and Emilsson, G.				
TITLE		Nucleozymes with endonuclease activity				
JOURNAL		Patent: WO 0159102-A 108 16-AUG-2001;				
		RIBOZYME PHARMACEUTICALS, INC. (US) ; Yale University (US)				


```
/note="2'-O-Methyl"
misc_feature 22..27
/note="2'-O-Methyl"
28
misc_feature 11 a 3 c 9 g 4 t 1 others
BASE COUNT
ORIGIN
Query Match 100.0%; Score 15; DB 6; Length 28;
Best Local Similarity 80.0%; Pred. No. 9.3e+02;
Matches 12; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGAUAACGUGAAGAU 15
Db 7 AGATAACGTGAAGAT 21

RESULT 6
AX214241
LOCUS AX214241 28 bp mRNA linear PAT 06-SEP-2001
DEFINITION Sequence 54 from Patent WO0159102.
ACCESSION AX214241
VERSION AX214241.1 GI:15524318
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct
artificial sequences.
REFERENCE 1 (bases 1 to 28)
AUTHORS Breaker,R. and Emilsson,G.
TITLE Nucleozymes with endonuclease activity
JOURNAL Patent: WO 0159102-A 54 16-AUG-2001;
RIBOZYME PHARMACEUTICALS, INC. (US) ; Yale University (US)
FEATURES
source 1..28
/organism="synthetic construct"
/db_xref="taxon:32630"
/note="Nucleic Acid"
misc_feature 1..5
/note="2'-O-Methyl"
23..27
misc_feature 28
/note="2'-O-Methyl"
28
BASE COUNT 11 a 3 c 9 g 4 t 1 others
ORIGIN
Query Match 100.0%; Score 15; DB 6; Length 28;
Best Local Similarity 80.0%; Pred. No. 9.3e+02;
Matches 12; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGAUAACGUGAAGAU 15
Db 7 AGATAACGTGAAGAT 21

RESULT 7
AX214242
LOCUS AX214242 28 bp mRNA linear PAT 06-SEP-2001
DEFINITION Sequence 55 from Patent WO0159102.
ACCESSION AX214242
VERSION AX214242.1 GI:15524319
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct
artificial sequences.
REFERENCE 1 (bases 1 to 28)
AUTHORS Breaker,R. and Emilsson,G.
TITLE Nucleozymes with endonuclease activity
JOURNAL Patent: WO 0159102-A 55 16-AUG-2001;
RIBOZYME PHARMACEUTICALS, INC. (US) ; Yale University (US)
FEATURES
source 1..28
/organism="synthetic construct"
```

```
/db_xref="taxon:32630"
/note="Nucleic Acid"
1..4
misc_feature 11 a 3 c 9 g 4 t 1 others
/note="2'-O-Methyl"
24..27
misc_feature 28
/note="2'-O-Methyl"
28
BASE COUNT 11 a 3 c 9 g 4 t 1 others
ORIGIN
Query Match 100.0%; Score 15; DB 6; Length 28;
Best Local Similarity 80.0%; Pred. No. 9.3e+02;
Matches 12; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGAUAACGUGAAGAU 15
Db 7 AGATAACGTGAAGAT 21

RESULT 8
AX214243
LOCUS AX214243 28 bp mRNA linear PAT 06-SEP-2001
DEFINITION Sequence 56 from Patent WO0159102.
ACCESSION AX214243
VERSION AX214243.1 GI:15524320
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct
artificial sequences.
REFERENCE 1 (bases 1 to 28)
AUTHORS Breaker,R. and Emilsson,G.
TITLE Nucleozymes with endonuclease activity
JOURNAL Patent: WO 0159102-A 56 16-AUG-2001;
RIBOZYME PHARMACEUTICALS, INC. (US) ; Yale University (US)
FEATURES
source 1..28
/organism="synthetic construct"
/db_xref="taxon:32630"
/note="Nucleic Acid"
1..4
misc_feature 23..27
/note="2'-O-Methyl"
28
BASE COUNT 11 a 3 c 9 g 4 t 1 others
ORIGIN
Query Match 100.0%; Score 15; DB 6; Length 28;
Best Local Similarity 80.0%; Pred. No. 9.3e+02;
Matches 12; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGAUAACGUGAAGAU 15
Db 7 AGATAACGTGAAGAT 21

RESULT 9
AX214244
LOCUS AX214244 28 bp mRNA linear PAT 06-SEP-2001
DEFINITION Sequence 57 from Patent WO0159102.
ACCESSION AX214244
VERSION AX214244.1 GI:15524321
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct
artificial sequences.
REFERENCE 1 (bases 1 to 28)
AUTHORS Breaker,R. and Emilsson,G.
TITLE Nucleozymes with endonuclease activity
JOURNAL Patent: WO 0159102-A 57 16-AUG-2001;
RIBOZYME PHARMACEUTICALS, INC. (US) ; Yale University (US)
```

FEATURES
source Location/Qualifiers
1..28
/organism="synthetic construct"
/db_xref="taxon:32630"
/note="Nucleic Acid"
1..6
/note="2'-O-Methyl"
20..27
/note="2'-O-Methyl"
28
/note="n stands for inverted deoxyabasic derivative"
11 a 3 c 9 g 4 t 1 others
ORIGIN
Query Match 100.0%; Score 15; DB 6; Length 28;
Best Local Similarity 80.0%; Pred. No. 9.3e+02;
Matches 12; Conservative 3; Mismatches 0; Indels 0; Gaps 0;
Qy 1 AGAUACGUGAAGAU 15
||||:||||:||||:
Db 7 AGATAACGTGAAGAT 21
RESULT 10
LOCUS AX214245 28 bp mRNA linear PAT 06-SEP-2001
DEFINITION Sequence 58 from Patent WO0159102.
ACCESSION AX214245
VERSION AX214245.1 GI:15524322
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct.
artificial sequences.
REFERENCE 1 (bases 1 to 28)
AUTHORS Breaker, R. and Emilsson, G.
TITLE Nucleozymes with endonuclease activity
JOURNAL Patent: WO 0159102-A 58 16-AUG-2001;
RIBOZYME PHARMACEUTICALS, INC. (US) ; Yale University (US)
FEATURES
source Location/Qualifiers
1..28
/organism="synthetic construct"
/db_xref="taxon:32630"
/note="Nucleic Acid"
1..6
/note="2'-O-Methyl"
12
/note="2'-O-Methyl"
21..27
/note="2'-O-Methyl"
28
/note="n stands for inverted deoxyabasic derivative"
11 a 3 c 9 g 4 t 1 others
ORIGIN
Query Match 100.0%; Score 15; DB 6; Length 28;
Best Local Similarity 80.0%; Pred. No. 9.3e+02;
Matches 12; Conservative 3; Mismatches 0; Indels 0; Gaps 0;
Qy 1 AGAUACGUGAAGAU 15
||||:||||:||||:
Db 7 AGATAACGTGAAGAT 21
RESULT 11
LOCUS AX214246 28 bp mRNA linear PAT 06-SEP-2001
DEFINITION Sequence 59 from Patent WO0159102.
ACCESSION AX214246
VERSION AX214246.1 GI:15524323
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct.
artificial sequences..

REFERENCE 1 (bases 1 to 28)
AUTHORS Breaker, R. and Emilsson, G.
TITLE Nucleozymes with endonuclease activity
JOURNAL Patent: WO 0159102-A 59 16-AUG-2001;
RIBOZYME PHARMACEUTICALS, INC. (US) ; Yale University (US)
FEATURES
source Location/Qualifiers
1..28
/organism="synthetic construct"
/db_xref="taxon:32630"
/note="Nucleic Acid"
1..7
/note="2'-O-Methyl"
21..27
/note="2'-O-Methyl"
28
/note="n stands for inverted deoxyabasic derivative"
11 a 3 c 9 g 4 t 1 others
ORIGIN
Query Match 100.0%; Score 15; DB 6; Length 28;
Best Local Similarity 80.0%; Pred. No. 9.3e+02;
Matches 12; Conservative 3; Mismatches 0; Indels 0; Gaps 0;
Qy 1 AGAUACGUGAAGAU 15
||||:||||:||||:
Db 7 AGATAACGTGAAGAT 21
RESULT 12
LOCUS AX214247 28 bp mRNA linear PAT 06-SEP-2001
DEFINITION Sequence 60 from Patent WO0159102.
ACCESSION AX214247
VERSION AX214247.1 GI:15524324
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct.
artificial sequences.
REFERENCE 1 (bases 1 to 28)
AUTHORS Breaker, R. and Emilsson, G.
TITLE Nucleozymes with endonuclease activity
JOURNAL Patent: WO 0159102-A 60 16-AUG-2001;
RIBOZYME PHARMACEUTICALS, INC. (US) ; Yale University (US)
FEATURES
source Location/Qualifiers
1..28
/organism="synthetic construct"
/db_xref="taxon:32630"
/note="Nucleic Acid"
1..6
/note="2'-O-Methyl"
21..27
/note="2'-O-Methyl"
28
/note="n stands for inverted deoxyabasic derivative"
11 a 3 c 9 g 4 t 1 others
ORIGIN
Query Match 100.0%; Score 15; DB 6; Length 28;
Best Local Similarity 80.0%; Pred. No. 9.3e+02;
Matches 12; Conservative 3; Mismatches 0; Indels 0; Gaps 0;
Qy 1 AGAUACGUGAAGAU 15
||||:||||:||||:
Db 7 AGATAACGTGAAGAT 21
RESULT 13
LOCUS AX214248 28 bp mRNA linear PAT 06-SEP-2001
DEFINITION Sequence 61 from Patent WO0159102.
ACCESSION AX214248
VERSION AX214248 GI:15524325
KEYWORDS

```

SOURCE          synthetic construct.
ORGANISM         synthetic construct
REFERENCE        1 (bases 1 to 28)
AUTHORS         Breaker,R. and Emilsson,G.
TITLE           Nucleozymes with endonuclease activity
JOURNAL         Patent: WO 0159102-A 61 16-AUG-2001;
                RIBOZYME PHARMACEUTICALS, INC. (US) ; Yale University (US)
FEATURES
  source        1..28
                /organism="synthetic construct"
                /db_xref="taxon:32630"
                /note="Nucleic Acid"
  misc_feature  1..6
                /note="2'-O-Methyl"
  misc_feature  18
                /note="2'-O-Methyl"
  misc_feature  21..27
                /note="2'-O-Methyl"
  misc_feature  28
                /note="2'-O-Methyl"
  BASE COUNT    11 a 3 c 9 g 4 t 1 others
  ORIGIN
Query Match     100.0%; Score 15; DB 6; Length 28;
Best Local Similarity 80.0%; Pred. No. 9.3e+02;
Matches 12; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGAUAACGUGAAGAU 15
Db 7 AGATAACGTGAAGAT 21

RESULT 14
AX214249
LOCUS           AX214249                28 bp      mRNA      linear      PAT 06-SEP-2001
DEFINITION     Sequence 62 from Patent WO0159102.
ACCESSION      AX214249
VERSION        AX214249.1 GI:15524326
KEYWORDS
SOURCE         synthetic construct.
ORGANISM       synthetic construct
REFERENCE      1 (bases 1 to 28)
AUTHORS        Breaker,R. and Emilsson,G.
TITLE          Nucleozymes with endonuclease activity
JOURNAL        Patent: WO 0159102-A 62 16-AUG-2001;
                RIBOZYME PHARMACEUTICALS, INC. (US) ; Yale University (US)
FEATURES
  source        1..28
                /organism="synthetic construct"
                /db_xref="taxon:32630"
                /note="Nucleic Acid"
  misc_feature  1..6
                /note="2'-O-Methyl"
  misc_feature  17
                /note="2'-O-Methyl"
  misc_feature  21..27
                /note="2'-O-Methyl"
  misc_feature  28
                /note="2'-O-Methyl"
  BASE COUNT    11 a 3 c 9 g 4 t 1 others
  ORIGIN
Query Match     100.0%; Score 15; DB 6; Length 28;
Best Local Similarity 80.0%; Pred. No. 9.3e+02;
Matches 12; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGAUAACGUGAAGAU 15
Db 7 AGATAACGTGAAGAT 21

```

```

RESULT 15
AX214250
LOCUS           AX214250                28 bp      mRNA      linear      PAT 06-SEP-2001
DEFINITION     Sequence 63 from Patent WO0159102.
ACCESSION      AX214250
VERSION        AX214250.1 GI:15524327
KEYWORDS
SOURCE         synthetic construct.
ORGANISM       synthetic construct
REFERENCE      1 (bases 1 to 28)
AUTHORS        Breaker,R. and Emilsson,G.
TITLE          Nucleozymes with endonuclease activity
JOURNAL        Patent: WO 0159102-A 63 16-AUG-2001;
                RIBOZYME PHARMACEUTICALS, INC. (US) ; Yale University (US)
FEATURES
  source        1..28
                /organism="synthetic construct"
                /db_xref="taxon:32630"
                /note="Nucleic Acid"
  misc_feature  1..6
                /note="2'-O-Methyl"
  misc_feature  11
                /note="2'-O-Methyl"
  misc_feature  16
                /note="2'-O-Methyl"
  misc_feature  21..27
                /note="2'-O-Methyl"
  misc_feature  28
                /note="2'-O-Methyl"
  BASE COUNT    11 a 3 c 9 g 4 t 1 others
  ORIGIN
Query Match     100.0%; Score 15; DB 6; Length 28;
Best Local Similarity 80.0%; Pred. No. 9.3e+02;
Matches 12; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGAUAACGUGAAGAU 15
Db 7 AGATAACGTGAAGAT 21

Search completed: July 6, 2003, 14:51:11
Job time : 493.182 secs

```

GenCore version 5.1.6
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: July 6, 2003, 14:25:15 ; Search time 160.909 Seconds

(without alignments)
209.932 Million cell updates/sec

Title: US-09-780-929-97

Perfect score: 15

Sequence: 1 agaaacgugaagau 15

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 2185239 seqs, 1125999159 residues

Total number of hits satisfying chosen parameters: 2274872

Minimum DB seq length: 0

Maximum DB seq length: 60

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

N_Geneseq_101002.*

1: /SID22/gcgdata/geneseq/geneseq-emb1/NA1980.DAT.*
2: /SID22/gcgdata/geneseq/geneseq-emb1/NA1981.DAT.*
3: /SID22/gcgdata/geneseq/geneseq-emb1/NA1982.DAT.*
4: /SID22/gcgdata/geneseq/geneseq-emb1/NA1983.DAT.*
5: /SID22/gcgdata/geneseq/geneseq-emb1/NA1984.DAT.*
6: /SID22/gcgdata/geneseq/geneseq-emb1/NA1985.DAT.*
7: /SID22/gcgdata/geneseq/geneseq-emb1/NA1986.DAT.*
8: /SID22/gcgdata/geneseq/geneseq-emb1/NA1987.DAT.*
9: /SID22/gcgdata/geneseq/geneseq-emb1/NA1988.DAT.*
10: /SID22/gcgdata/geneseq/geneseq-emb1/NA1989.DAT.*
11: /SID22/gcgdata/geneseq/geneseq-emb1/NA1990.DAT.*
12: /SID22/gcgdata/geneseq/geneseq-emb1/NA1991.DAT.*
13: /SID22/gcgdata/geneseq/geneseq-emb1/NA1992.DAT.*
14: /SID22/gcgdata/geneseq/geneseq-emb1/NA1993.DAT.*
15: /SID22/gcgdata/geneseq/geneseq-emb1/NA1994.DAT.*
16: /SID22/gcgdata/geneseq/geneseq-emb1/NA1995.DAT.*
17: /SID22/gcgdata/geneseq/geneseq-emb1/NA1996.DAT.*
18: /SID22/gcgdata/geneseq/geneseq-emb1/NA1997.DAT.*
19: /SID22/gcgdata/geneseq/geneseq-emb1/NA1998.DAT.*
20: /SID22/gcgdata/geneseq/geneseq-emb1/NA1999.DAT.*
21: /SID22/gcgdata/geneseq/geneseq-emb1/NA2000.DAT.*
22: /SID22/gcgdata/geneseq/geneseq-emb1/NA2001A.DAT.*
23: /SID22/gcgdata/geneseq/geneseq-emb1/NA2001B.DAT.*
24: /SID22/gcgdata/geneseq/geneseq-emb1/NA2002.DAT.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	15	100.0	15	22	AA12347
2	15	100.0	27	22	AA12347
3	15	100.0	27	22	AA12295
4	15	100.0	27	22	AA12296
5	15	100.0	27	22	AA12297
6	15	100.0	27	22	AA12298
7	15	100.0	27	22	AA12299
8	15	100.0	27	22	AA12300
9	15	100.0	27	22	AA12301
	15	100.0	27	22	AA12302

10	15	100.0	27	22	AA12303	DNA encoding class
11	15	100.0	27	22	AA12304	DNA encoding class
12	15	100.0	27	22	AA12305	DNA encoding class
13	15	100.0	27	22	AA12306	DNA encoding class
14	15	100.0	27	22	AA12307	DNA encoding class
15	15	100.0	27	22	AA12308	DNA encoding class
16	15	100.0	27	22	AA12309	DNA encoding class
17	15	100.0	27	22	AA12310	DNA encoding class
18	15	100.0	27	22	AA12311	DNA encoding class
19	15	100.0	27	22	AA12312	DNA encoding class
20	15	100.0	27	22	AA12313	DNA encoding class
21	15	100.0	27	22	AA12314	DNA encoding class
22	15	100.0	27	22	AA12315	DNA encoding class
23	15	100.0	27	22	AA12316	DNA encoding class
24	15	100.0	27	22	AA12317	DNA encoding class
25	15	100.0	27	22	AA12318	DNA encoding class
26	15	100.0	27	22	AA12319	DNA encoding class
27	15	100.0	27	22	AA12320	DNA encoding class
28	15	100.0	27	22	AA12321	DNA encoding class
29	15	100.0	27	22	AA12322	DNA encoding class
30	15	100.0	27	22	AA12323	DNA encoding class
31	15	100.0	27	22	AA12324	DNA encoding class
32	15	100.0	27	22	AA12325	DNA encoding class
33	15	100.0	27	22	AA12326	DNA encoding class
34	15	100.0	27	22	AA12327	DNA encoding class
35	15	100.0	27	22	AA12328	DNA encoding class
36	15	100.0	27	22	AA12329	DNA encoding class
37	15	100.0	27	22	AA12330	DNA encoding class
38	15	100.0	27	22	AA12331	DNA encoding class
39	15	100.0	27	22	AA12332	DNA encoding class
40	15	100.0	27	22	AA12333	DNA encoding class
41	15	100.0	27	22	AA12334	DNA encoding class
42	15	100.0	27	22	AA12335	DNA encoding class
43	15	100.0	27	22	AA12336	DNA encoding class
44	15	100.0	27	22	AA12337	DNA encoding class
45	15	100.0	27	22	AA12404	DNA encoding class

ALIGNMENTS

RESULT 1

AA12347
ID AA12347 standard; DNA; 15 BP.

XX AA12347;

XX 21-NOV-2001 (first entry)

XX DNA encoding deoxyribozyme #7.

XX Deoxyribozyme; cytostatic; endonuclease; RNA cleavage; DNA cleavage;
KW gene therapy; plant; fungus; bacteria; mammal; ribozyme; ss.

XX Synthetic.

XX WO200159102-A2.

XX 16-AUG-2001.

XX 08-FEB-2001; 2001WO-US04223.

XX 08-FEB-2000; 2000US-0181360.

XX 31-MAR-2000; 2000US-0193646.

XX (RIBO-) RIBOZYME PHARM INC.

XX (UYFA) UNIV YALE.

XX Breaker R, Beigelman L, Emilsson G;

XX WPI; 2001-536526/59.

XX New nucleic acids with endonuclease activity, such as ribozymes and

PT nucleozymes, for modulating gene expression in a plant, mammalian,
XX bacterial or fungal cell
PS Claim 49; Page 77; 96pp; English.
XX
CC The invention relates to nucleic acid molecules with endonuclease
CC activity, which are particularly useful for cleavage of RNA or DNA.
CC The nucleic acids are used in a pharmaceutical composition and are used
CC to modulate expression of a gene in a plant, mammalian, bacterial or
CC fungal cell. They are used to cleave a separate nucleic acid, preferably
CC RNA. The nucleic acids are used to inhibit gene expression and/or cell
CC proliferation, and can be used to treat a disease or condition. More
CC than one nucleic acid can be independently targeted to the same or
CC different sites in a cell. The nucleic acids may be used to study DNA.
CC The modifications to the nucleic acids optimises their catalytic activity
CC and can maintain or enhance their activity. They exhibit a high degree
CC of specificity for RNA. The present sequence represents the coding
CC sequence of deoxyribozyme #7 used in the method of the invention.
XX
SQ Sequence 15 BP; 7 A; 1 C; 4 G; 3 U; 0 other;
Query Match 100.0%; Score 15; DB 22; Length 15;
Best Local Similarity 100.0%; Pred. No. 71;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 AGAUAACGUGAAGAU 15
|||||
Db 1 AGAUAACGUGAAGAU 15
RESULT 2
AAS12295
ID AAS12295 standard; DNA; 27 BP.
AC AAS12295;
XX
XX 21-NOV-2001 (first entry)
XX
XX DNA encoding class V ribozyme #7.
XX
XX Ribozyme; cytosstatic; endonuclease; RNA cleavage; DNA cleavage;
KW gene therapy; plant; fungus; bacteria; mammal; ss.
XX
XX Synthetic.
XX
XX Key Location/Qualifiers
FH modified_base 1..4
FT /*tag= a
FT /mod_base= OTHER
FT /note= "OTHER = 2'-O-methyl nucleotides"
FT modified_base 23..27
FT /*tag= b
FT /mod_base= OTHER
FT /note= "OTHER = 2'-O-methyl nucleotides"
FT modified_base 27
FT /*tag= c
FT /mod_base= c
FT /note= "3',3'-inverted deoxybasic moiety"
XX
XX WO200159102-A2.
XX
XX 16-AUG-2001.
XX
XX 08-FEB-2001; 2001WO-US04223.
XX
XX 08-FEB-2000; 2000US-0181360.
PR 31-MAR-2000; 2000US-0193646.
XX
XX (RIBO-) RIBOZYME PHARM INC.
PA (UYIA) UNIV YALE.
XX
XX Breaker R, Beigelman L, Emilsson G;
PI
XX

DR WPI; 2001-536526/59.
XX
XX New nucleic acids with endonuclease activity, such as ribozymes and
PT nucleozymes, for modulating gene expression in a plant, mammalian,
XX bacterial or fungal cell
PS Example 1; Page 71; 96pp; English.
XX
CC The invention relates to nucleic acid molecules with endonuclease
CC activity, which are particularly useful for cleavage of RNA or DNA.
CC The nucleic acids are used in a pharmaceutical composition and are used
CC to modulate expression of a gene in a plant, mammalian, bacterial or
CC fungal cell. They are used to cleave a separate nucleic acid, preferably
CC RNA. The nucleic acids are used to inhibit gene expression and/or cell
CC proliferation, and can be used to treat a disease or condition. More
CC than one nucleic acid can be independently targeted to the same or
CC different sites in a cell. The nucleic acids may be used to study DNA.
CC The modifications to the nucleic acids optimises their catalytic activity
CC and can maintain or enhance their activity. They exhibit a high degree
CC of specificity for RNA. The present sequence represents the coding
CC sequence of class V ribozyme #7 used in the method of the invention.
XX
SQ Sequence 27 BP; 11 A; 3 C; 9 G; 4 U; 0 other;
Query Match 100.0%; Score 15; DB 22; Length 27;
Best Local Similarity 100.0%; Pred. No. 74;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 AGAUAACGUGAAGAU 15
|||||
Db 7 AGAUAACGUGAAGAU 21
RESULT 3
AAS12296
ID AAS12296 standard; DNA; 27 BP.
XX
XX AAS12296;
XX
XX 21-NOV-2001 (first entry)
XX
XX DNA encoding class V ribozyme #8.
XX
XX Ribozyme; cytosstatic; endonuclease; RNA cleavage; DNA cleavage;
KW gene therapy; plant; fungus; bacteria; mammal; ss.
XX
XX Synthetic.
XX
XX Key Location/Qualifiers
FH modified_base 1..6
FT /*tag= a
FT /mod_base= OTHER
FT /note= "OTHER = 2'-O-methyl nucleotides"
FT modified_base 20..27
FT /*tag= b
FT /mod_base= OTHER
FT /note= "OTHER = 2'-O-methyl nucleotides"
FT modified_base 27
FT /*tag= c
FT /mod_base= c
FT /note= "3',3'-inverted deoxybasic moiety"
XX
XX WO200159102-A2.
XX
XX 16-AUG-2001.
XX
XX 08-FEB-2001; 2001WO-US04223.
XX
XX 08-FEB-2000; 2000US-0181360.
PR 31-MAR-2000; 2000US-0193646.
XX
XX (RIBO-) RIBOZYME PHARM INC.
PA (UYIA) UNIV YALE.
XX
XX


```
XX PI Breaker R, Beigelman L, Emilsson G;
XX PA WPI; 2001-536526/59.
XX DR
XX XX
XX PT New nucleic acids with endonuclease activity, such as ribozymes and
XX FT nucleozymes, for modulating gene expression in a plant, mammalian,
XX FT bacterial or fungal cell
XX PS
XX PS Example 1; Page 71; 96pp; English.
XX CC The invention relates to nucleic acid molecules with endonuclease
XX CC activity, which are particularly useful for cleavage of RNA or DNA.
XX CC The nucleic acids are used in a pharmaceutical composition and are used
XX CC to modulate expression of a gene in a plant, mammalian, bacterial or
XX CC fungal cell. They are used to cleave a separate nucleic acid, preferably
XX CC RNA. The nucleic acids are used to inhibit gene expression and/or cell
XX CC proliferation, and can be used to treat a disease or condition. More
XX CC than one nucleic acid can be independently targeted to the same or
XX CC different sites in a cell. The nucleic acids may be used to study DNA.
XX CC The modifications to the nucleic acids optimises their catalytic activity
XX CC and can maintain or enhance their activity. They exhibit a high degree
XX CC of specificity for RNA. The present sequence represents the coding
XX CC sequence of class V ribozyme #8 used in the method of the invention.
XX SQ Sequence 27 BP; 11 A; 3 C; 9 G; 4 U; 0 other;
Query Match 100.0%; Score 15; DB 22; Length 27;
Best Local Similarity 100.0%; Pred. No. 74;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 AGAUAACGUGAAGAU 15
Db 7 AGAUAACGUGAAGAU 21
RESULT 4
AAS12297
ID AAS12297 standard; DNA; 27 BP.
XX AC AAS12297;
XX DT 21-NOV-2001 (first entry)
XX DE DNA encoding class V ribozyme #9.
XX KW Ribozyme; cytostatic; endonuclease; RNA cleavage; DNA cleavage;
XX KW gene therapy; plant; fungus; bacteria; mammal; ss.
XX OS Synthetic.
XX FH Key Location/Qualifiers
XX FT modified_base 1..6
XX FT /tag= a
XX FT /mod_base= OTHER
XX FT /note= "OTHER = 2'-O-methyl nucleotides"
XX FT modified_base 21..27
XX FT /tag= b
XX FT /mod_base= OTHER
XX FT /note= "OTHER = 2'-O-methyl nucleotides"
XX FT modified_base 27
XX FT /tag= c
XX FT /mod_base= c
XX FT /note= "3',3'-inverted deoxyabasic moiety"
XX PN WO200159102-A2.
XX PD 16-AUG-2001.
XX PF 08-FEB-2001; 2001WO-US04223.
XX PR 08-FEB-2000; 2000US-0181360.
XX PR 31-MAR-2000; 2000US-0193646.
```

```
XX XX (RIBO-) RIBOZYME PHARM INC.
XX PA (UYVA ) UNIV YALE.
XX XX
XX PI Breaker R, Beigelman L, Emilsson G;
XX DR WPI; 2001-536526/59.
XX XX
XX PT New nucleic acids with endonuclease activity, such as ribozymes and
XX FT nucleozymes, for modulating gene expression in a plant, mammalian,
XX FT bacterial or fungal cell
XX PS
XX PS Example 1; Page 71; 96pp; English.
XX CC The invention relates to nucleic acid molecules with endonuclease
XX CC activity, which are particularly useful for cleavage of RNA or DNA.
XX CC The nucleic acids are used in a pharmaceutical composition and are used
XX CC to modulate expression of a gene in a plant, mammalian, bacterial or
XX CC fungal cell. They are used to cleave a separate nucleic acid, preferably
XX CC RNA. The nucleic acids are used to inhibit gene expression and/or cell
XX CC proliferation, and can be used to treat a disease or condition. More
XX CC than one nucleic acid can be independently targeted to the same or
XX CC different sites in a cell. The nucleic acids may be used to study DNA.
XX CC The modifications to the nucleic acids optimises their catalytic activity
XX CC and can maintain or enhance their activity. They exhibit a high degree
XX CC of specificity for RNA. The present sequence represents the coding
XX CC sequence of class V ribozyme #9 used in the method of the invention.
XX SQ Sequence 27 BP; 11 A; 3 C; 9 G; 4 U; 0 other;
Query Match 100.0%; Score 15; DB 22; Length 27;
Best Local Similarity 100.0%; Pred. No. 74;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 AGAUAACGUGAAGAU 15
Db 7 AGAUAACGUGAAGAU 21
RESULT 5
AAS12298
ID AAS12298 standard; DNA; 27 BP.
XX AC AAS12298;
XX DT 21-NOV-2001 (first entry)
XX DE DNA encoding class V ribozyme #10.
XX KW Ribozyme; cytostatic; endonuclease; RNA cleavage; DNA cleavage;
XX KW gene therapy; plant; fungus; bacteria; mammal; ss.
XX OS Synthetic.
XX FH Key Location/Qualifiers
XX FT modified_base 1..6
XX FT /tag= a
XX FT /mod_base= OTHER
XX FT /note= "OTHER = 2'-O-methyl nucleotides"
XX FT modified_base 12
XX FT /tag= b
XX FT /mod_base= OTHER
XX FT /note= "OTHER = 2'-O-methyl nucleotide"
XX FT modified_base 21..27
XX FT /tag= c
XX FT /mod_base= OTHER
XX FT /note= "OTHER = 2'-O-methyl nucleotides"
XX FT modified_base 27
XX FT /tag= d
XX FT /mod_base= c
XX FT /note= "3',3'-inverted deoxyabasic moiety"
XX PN WO200159102-A2.
```

```
XX PD 16-AUG-2001.
XX XX
XX PF 08-FEB-2001; 2001WO-US04223.
XX XX
XX PR 08-FEB-2000; 2000US-0181360.
XX PR 31-MAR-2000; 2000US-0193646.
XX XX
XX PA (RIBO-) RIBOZYME PHARM INC.
XX PA (UYIA ) UNIV YALE.
XX XX
XX PI Breaker R, Beigelman L, Emilsson G;
XX XX WPI; 2001-536526/59.
XX DR
XX XX New nucleic acids with endonuclease activity, such as ribozymes and
XX PT nucleozymes, for modulating gene expression in a plant, mammalian,
XX PT bacterial or fungal cell -
XX XX
XX PS Example 1; Page 71; 96pp; English.
XX CC The invention relates to nucleic acid molecules with endonuclease
XX CC activity, which are particularly useful for cleavage of RNA or DNA.
XX CC The nucleic acids are used in a pharmaceutical composition and are used
XX CC to modulate expression of a gene in a plant, mammalian, bacterial or
XX CC fungal cell. They are used to cleave a separate nucleic acid, preferably
XX CC RNA. The nucleic acids are used to inhibit gene expression and/or cell
XX CC proliferation, and can be used to treat a disease or condition. More
XX CC than one nucleic acid can be independently targeted to the same or
XX CC different sites in a cell. The nucleic acids may be used to study DNA.
XX CC The modifications to the nucleic acids optimises their catalytic activity
XX CC and can maintain or enhance their activity. They exhibit a high degree
XX CC of specificity for RNA. The present sequence represents the coding
XX CC sequence of class V ribozyme #10 used in the method of the invention.
XX XX
XX SQ Sequence 27 BP; 11 A; 3 C; 9 G; 4 U; 0 other;
XX
Query Match 100.0%; Score 15; DB 22; Length 27;
Best Local Similarity 100.0%; Pred. No. 74;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGAUACGUGAAGAU 15
Db 7 AGAUACGUGAAGAU 21

RESULT 6
AAS12299
ID AAS12299 standard; DNA; 27 BP.
XX AC AAS12299;
XX DT 21-NOV-2001 (first entry)
XX DE DNA encoding class V ribozyme #11.
XX KW Ribozyme; cytosolic; endonuclease; RNA cleavage; DNA cleavage;
XX KW gene therapy; plant; fungus; bacteria; mammal; ss.
XX OS Synthetic.
XX FH Key Location/Qualifiers
XX FT modified_base 1..7
XX FT /tag= a
XX FT /mod_base= OTHER
XX FT /note= "OTHER = 2'-O-methyl nucleotides"
XX FT modified_base 21..27
XX FT /tag= b
XX FT /mod_base= OTHER
XX FT /note= "OTHER = 2'-O-methyl nucleotides"
XX FT modified_base 27
XX FT /tag= c
XX FT /mod_base= c
```

```
FT XX /note= "3',3'-inverted deoxyabasic moiety"
XX PN WO200159102-A2.
XX XX
XX PD 16-AUG-2001.
XX XX
XX PF 08-FEB-2001; 2001WO-US04223.
XX XX
XX PR 08-FEB-2000; 2000US-0181360.
XX PR 31-MAR-2000; 2000US-0193646.
XX XX
XX XX (RIBO-) RIBOZYME PHARM INC.
XX PA (UYIA ) UNIV YALE.
XX XX
XX PI Breaker R, Beigelman L, Emilsson G;
XX XX WPI; 2001-536526/59.
XX DR
XX XX New nucleic acids with endonuclease activity, such as ribozymes and
XX PT nucleozymes, for modulating gene expression in a plant, mammalian,
XX PT bacterial or fungal cell -
XX XX
XX PS Example 1; Page 71; 96pp; English.
XX CC The invention relates to nucleic acid molecules with endonuclease
XX CC activity, which are particularly useful for cleavage of RNA or DNA.
XX CC The nucleic acids are used in a pharmaceutical composition and are used
XX CC to modulate expression of a gene in a plant, mammalian, bacterial or
XX CC fungal cell. They are used to cleave a separate nucleic acid, preferably
XX CC RNA. The nucleic acids are used to inhibit gene expression and/or cell
XX CC proliferation, and can be used to treat a disease or condition. More
XX CC than one nucleic acid can be independently targeted to the same or
XX CC different sites in a cell. The nucleic acids may be used to study DNA.
XX CC The modifications to the nucleic acids optimises their catalytic activity
XX CC and can maintain or enhance their activity. They exhibit a high degree
XX CC of specificity for RNA. The present sequence represents the coding
XX CC sequence of class V ribozyme #11 used in the method of the invention.
XX XX
XX SQ Sequence 27 BP; 11 A; 3 C; 9 G; 4 U; 0 other;
XX
Query Match 100.0%; Score 15; DB 22; Length 27;
Best Local Similarity 100.0%; Pred. No. 74;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGAUACGUGAAGAU 15
Db 7 AGAUACGUGAAGAU 21

RESULT 7
AAS12300
ID AAS12300 standard; DNA; 27 BP.
XX AC AAS12300;
XX DT 21-NOV-2001 (first entry)
XX DE DNA encoding class V ribozyme #12.
XX KW Ribozyme; cytosolic; endonuclease; RNA cleavage; DNA cleavage;
XX KW gene therapy; plant; fungus; bacteria; mammal; ss.
XX OS Synthetic.
XX FH Key Location/Qualifiers
XX FT modified_base 1..6
XX FT /tag= a
XX FT /mod_base= OTHER
XX FT /note= "OTHER = 2'-O-methyl nucleotides"
XX FT modified_base 21..27
XX FT /tag= b
XX FT /mod_base= OTHER
XX FT /note= "OTHER = 2'-O-methyl nucleotides"
```

```

FT modified_base 27 /*tag= c
FT /*mod_base= c
FT /*note= "3',3'-inverted deoxyabasic moiety"
XX
PN WO200159102-A2.
XX
PD 16-AUG-2001.
XX
PF 08-FEB-2001; 2001WO-US04223.
XX
PR 08-FEB-2000; 2000US-0181360.
PR 31-MAR-2000; 2000US-0193646.
XX
PA (RIBO-) RIBOZYME PHARM INC.
PA (UYVA ) UNIV YALE.
XX
PI Breaker R, Beigelman L, Emilsson G;
XX
DR WPI; 2001-536526/59.
XX
PS New nucleic acids with endonuclease activity, such as ribozymes and
PT nucleozymes, for modulating gene expression in a plant, mammalian,
PT bacterial or fungal cell
XX
XX Example 1; Page 71; 96pp; English.
XX
CC The invention relates to nucleic acid molecules with endonuclease
CC activity, which are particularly useful for cleavage of RNA or DNA.
CC The nucleic acids are used in a pharmaceutical composition and are used
CC to modulate expression of a gene in a plant, mammalian, bacterial or
CC fungal cell. They are used to cleave a separate nucleic acid, preferably
CC RNA. The nucleic acids are used to inhibit gene expression and/or cell
CC proliferation, and can be used to treat a disease or condition. More
CC than one nucleic acid can be independently targeted to the same or
CC different sites in a cell. The nucleic acids may be used to study DNA.
CC The modifications to the nucleic acids optimises their catalytic activity
CC and can maintain or enhance their activity. They exhibit a high degree
CC of specificity for RNA. The present sequence represents the coding
CC sequence of class V ribozyme #12 used in the method of the invention.
XX
SQ Sequence 27 BP; 11 A; 3 C; 9 G; 4 U; 0 other;
Query Match 100.0%; Score 15; DB 22; Length 27;
Best Local Similarity 100.0%; Pred. No. 74;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 AGAUAACGUGAAGAU 15
Db |||||
7 AGAUAACGUGAAGAU 21
RESULT 8
AAS12301
ID AAS12301 standard; DNA; 27 BP.
XX
AC AAS12301;
XX
DT 21-NOV-2001 (first entry)
XX
DE DNA encoding class V ribozyme #13.
XX
KW Ribozyme; cytostatic; endonuclease; RNA cleavage; DNA cleavage;
KW gene therapy; plant; fungus; bacteria; mammal; ss.
XX
OS Synthetic.
XX
FH Key Location/Qualifiers
FT modified_base 1..6 /*tag= a
FT /*mod_base= OTHER
FT /*note= "2'-O-methyl nucleotides"
FT modified_base 18

```

```

FT /*tag= b
FT /*mod_base= a
FT /*note= "2'-O-methyl nucleotide"
FT modified_base 21..27
FT /*tag= c
FT /*mod_base= OTHER
FT /*note= "2'-O-methyl nucleotides"
FT modified_base 27
FT /*tag= d
FT /*mod_base= c
FT /*note= "3',3'-inverted deoxyabasic moiety"
XX
PN WO200159102-A2.
XX
PD 16-AUG-2001.
XX
PF 08-FEB-2001; 2001WO-US04223.
XX
PR 08-FEB-2000; 2000US-0181360.
PR 31-MAR-2000; 2000US-0193646.
XX
PA (RIBO-) RIBOZYME PHARM INC.
PA (UYVA ) UNIV YALE.
XX
PI Breaker R, Beigelman L, Emilsson G;
XX
DR WPI; 2001-536526/59.
XX
PS New nucleic acids with endonuclease activity, such as ribozymes and
PT nucleozymes, for modulating gene expression in a plant, mammalian,
PT bacterial or fungal cell
XX
XX Example 1; Page 71; 96pp; English.
XX
CC The invention relates to nucleic acid molecules with endonuclease
CC activity, which are particularly useful for cleavage of RNA or DNA.
CC The nucleic acids are used in a pharmaceutical composition and are used
CC to modulate expression of a gene in a plant, mammalian, bacterial or
CC fungal cell. They are used to cleave a separate nucleic acid, preferably
CC RNA. The nucleic acids are used to inhibit gene expression and/or cell
CC proliferation, and can be used to treat a disease or condition. More
CC than one nucleic acid can be independently targeted to the same or
CC different sites in a cell. The nucleic acids may be used to study DNA.
CC The modifications to the nucleic acids optimises their catalytic activity
CC and can maintain or enhance their activity. They exhibit a high degree
CC of specificity for RNA. The present sequence represents the coding
CC sequence of class V ribozyme #13 used in the method of the invention.
XX
SQ Sequence 27 BP; 11 A; 3 C; 9 G; 4 U; 0 other;
Query Match 100.0%; Score 15; DB 22; Length 27;
Best Local Similarity 100.0%; Pred. No. 74;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 AGAUAACGUGAAGAU 15
Db |||||
7 AGAUAACGUGAAGAU 21
RESULT 9
AAS12302
ID AAS12302 standard; DNA; 27 BP.
XX
AC AAS12302;
XX
DT 21-NOV-2001 (first entry)
XX
DE DNA encoding class V ribozyme #14.
XX
KW Ribozyme; cytostatic; endonuclease; RNA cleavage; DNA cleavage;
KW gene therapy; plant; fungus; bacteria; mammal; ss.
XX
OS Synthetic.

```

```
XX FH Key Location/Qualifiers
XX FT modified_base 1..6
XX FT /tag= a
XX FT /mod_base= OTHER
XX FT /note= "OTHER = 2'-O-methyl nucleotides"
XX FT modified_base 17
XX FT /tag= b
XX FT /mod_base= a
XX FT /note= "OTHER = 2'-O-methyl nucleotide"
XX FT modified_base 21..27
XX FT /tag= c
XX FT /mod_base= OTHER
XX FT /note= "OTHER = 2'-O-methyl nucleotides"
XX FT modified_base 27
XX FT /tag= d
XX FT /mod_base= c
XX FT /note= "3',3'-inverted deoxybasic moiety"
XX PN WO200159102-A2.
XX PD 16-AUG-2001.
XX PF 08-FEB-2001; 2001WO-US04223.
XX PR 08-FEB-2000; 2000US-0181360.
XX PR 31-MAR-2000; 2000US-0193646.
XX PA (RIBO-) RIBOZYME PHARM INC.
XX PA (UYVA ) UNIV YALE.
XX PI Breaker R, Beigelman L, Emilsson G;
XX XX WPI; 2001-536526/59.
XX XX New nucleic acids with endonuclease activity, such as ribozymes and
XX PT nucleozymes, for modulating gene expression in a plant, mammalian,
XX PT bacterial or fungal cell
XX PS Example 1; Page 71; 96pp; English.
XX CC The invention relates to nucleic acid molecules with endonuclease
XX CC activity, which are particularly useful for cleavage of RNA or DNA.
XX CC The nucleic acids are used in a pharmaceutical composition and are used
XX CC to modulate expression of a gene in a plant, mammalian, bacterial or
XX CC fungal cell. They are used to cleave a separate nucleic acid, preferably
XX CC RNA. The nucleic acids are used to inhibit gene expression and/or cell
XX CC proliferation, and can be used to treat a disease or condition. More
XX CC than one nucleic acid can be independently targeted to the same or
XX CC different sites in a cell. The nucleic acids may be used to study DNA.
XX CC The modifications to the nucleic acids optimises their catalytic activity
XX CC and can maintain or enhance their activity. They exhibit a high degree
XX CC of specificity for RNA. The present sequence represents the coding
XX CC sequence of class V ribozyme #14 used in the method of the invention.
XX SQ Sequence 27 BP; 11 A; 3 C; 9 G; 4 U; 0 other;
XX
XX Query Match 100.0%; Score 15; DB 22; Length 27;
XX Best Local Similarity 100.0%; Pred. No. 74;
XX Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX QY 1 AGAUAACGUGAAGAU 15
XX |||||
XX Db 7 AGAUAACGUGAAGAU 21
XX
XX RESULT 10
XX AAS12303
XX ID AAS12303 standard; DNA; 27 BP.
XX AC AAS12303;
XX XX
XX DT 21-NOV-2001 (first entry)
```

```
XX DE DNA encoding class V ribozyme #15.
XX KW Ribozyme; cytostatic; endonuclease; RNA cleavage; DNA cleavage;
XX KW gene therapy; plant; fungus; bacteria; mammal; ss.
XX OS Synthetic.
XX FH Key Location/Qualifiers
XX FT modified_base 1..6
XX FT /tag= a
XX FT /mod_base= OTHER
XX FT /note= "OTHER = 2'-O-methyl nucleotides"
XX FT modified_base 11
XX FT /tag= b
XX FT /mod_base= a
XX FT /note= "OTHER = 2'-O-methyl nucleotide"
XX FT modified_base 16
XX FT /tag= c
XX FT /mod_base= g
XX FT /note= "OTHER = 2'-O-methyl nucleotide"
XX FT modified_base 21..27
XX FT /tag= d
XX FT /mod_base= OTHER
XX FT /note= "OTHER = 2'-O-methyl nucleotides"
XX FT modified_base 27
XX FT /tag= e
XX FT /mod_base= c
XX FT /note= "3',3'-inverted deoxybasic moiety"
XX PN WO200159102-A2.
XX PD 16-AUG-2001.
XX PF 08-FEB-2001; 2001WO-US04223.
XX PR 08-FEB-2000; 2000US-0181360.
XX PR 31-MAR-2000; 2000US-0193646.
XX PA (RIBO-) RIBOZYME PHARM INC.
XX PA (UYVA ) UNIV YALE.
XX PI Breaker R, Beigelman L, Emilsson G;
XX XX WPI; 2001-536526/59.
XX XX New nucleic acids with endonuclease activity, such as ribozymes and
XX PT nucleozymes, for modulating gene expression in a plant, mammalian,
XX PT bacterial or fungal cell
XX PS Example 1; Page 71; 96pp; English.
XX CC The invention relates to nucleic acid molecules with endonuclease
XX CC activity, which are particularly useful for cleavage of RNA or DNA.
XX CC The nucleic acids are used in a pharmaceutical composition and are used
XX CC to modulate expression of a gene in a plant, mammalian, bacterial or
XX CC fungal cell. They are used to cleave a separate nucleic acid, preferably
XX CC RNA. The nucleic acids are used to inhibit gene expression and/or cell
XX CC proliferation, and can be used to treat a disease or condition. More
XX CC than one nucleic acid can be independently targeted to the same or
XX CC different sites in a cell. The nucleic acids may be used to study DNA.
XX CC The modifications to the nucleic acids optimises their catalytic activity
XX CC and can maintain or enhance their activity. They exhibit a high degree
XX CC of specificity for RNA. The present sequence represents the coding
XX CC sequence of class V ribozyme #15 used in the method of the invention.
XX SQ Sequence 27 BP; 11 A; 3 C; 9 G; 4 U; 0 other;
XX
XX Query Match 100.0%; Score 15; DB 22; Length 27;
XX Best Local Similarity 100.0%; Pred. No. 74;
XX Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX QY 1 AGAUAACGUGAAGAU 15
```

Db

|||||
7 AGAUAACGUGAGAU 21

RESULT 11
AAS12304

ID AAS12304 standard; DNA; 27 BP.
XX
XX AAS12304;
XX
XX 21-NOV-2001 (first entry)
XX
XX DNA encoding class V ribozyme #16.
XX
XX Ribozyme: cytosolic; endonuclease; RNA cleavage; DNA cleavage;
KW gene therapy; plant; fungus; bacteria; mammal; ss.
XX
XX Synthetic.

Key	Location/Qualifiers
modified_base 1..6	/*tag= a
FT FT	/mod_base= OTHER
FT FT	/note= "OTHER = 2'-O-methyl nucleotides"
modified_base 21..27	/*tag= b
FT FT	/mod_base= OTHER
FT FT	/note= "OTHER = 2'-O-methyl nucleotides"
modified_base 27	/*tag= c
FT FT	/mod_base= c
FT FT	/note= "3',3'-inverted deoxyabasic moiety"

WOZ001I59102-A2.
PN
XX
XX
XX
PD 16-AUG-2001.
XX
XX
XX 08-FEB-2001; 2001WO-US04223.
PF
XX
XX 08-FEB-2000; 2000US-0181360.
PR
XX 31-MAR-2000; 2000US-0193646.
XX
PA (RIBO-) RIBOZYME PHARM INC.
PA (UYIA) UNIV YALE.
XX
XX Breaker R, Beigelman L, Emilsson G;
PI
XX WPI; 2001-536526/59.
DR
XX
PT New nucleic acids with endonuclease activity, such as ribozymes and
PT nucleozymes, for modulating gene expression in a plant, mammalian,
PT bacterial or fungal cell
XX
XX Example 1; Page 71; 96pp; English.
PS
XX
CC The invention relates to nucleic acid molecules with endonuclease
CC activity, which are particularly useful for cleavage of RNA or DNA.
CC The nucleic acids are used in a pharmaceutical composition and are used
CC to modulate expression of a gene in a plant, mammalian, bacterial or
CC fungal cell. They are used to cleave a separate nucleic acid, preferably
CC RNA. The nucleic acids are used to inhibit gene expression and/or cell
CC proliferation, and can be used to treat a disease or condition. More
CC than one nucleic acid can be independently targeted to the same or
CC different sites in a cell. The nucleic acids may be used to study DNA.
CC The modifications to the nucleic acids optimises their catalytic activity
CC and can maintain or enhance their activity. They exhibit a high degree
CC of specificity for RNA. The present sequence represents the coding
CC sequence of class V ribozyme #16 used in the method of the invention.
XX
SQ Sequence 27 BP; 11 A; 3 C; 9 G; 4 U; 0 other;

Query Match 100.0%; Score 15; DB 22; Length 27;
Best Local Similarity 100.0%; Pred. No. 74;

	Matches	15;	Conservative	0;	Mismatches	0;	Indels	0;	Gaps	0;
Qy	1	AGAUACGUGAAGAU	15							
Db	7	AGAUACGUGAAGAU	21							
 RESULT 12										
AAS12305	ID	AAS12305 standard; DNA; 27 BP.								
XX	AC	AAS12305;								
XX	DT	21-NOV-2001 (first entry)								
XX	XX									
DE	DE	DNA encoding class V ribozyme #17.								
XX	XX									
KW	KW	Ribozyme; cytosstatic; endonuclease; RNA cleavage; DNA cleavage;								
KW	KW	gene therapy; plant; fungus; bacteria; mammal; ss.								
XX	OS	Synthetic..								
XX	XX									
FH	Key	Location/Qualifiers								
FT	modified_base	1..6								
FT	FT	/*tag= a								
FT	FT	/mod_base= OTHER								
FT	FT	/note= "OTHER = 2'-O-methyl nucleotides"								
FT	FT	19								
FT	FT	/*tag= b								
FT	FT	/mod_base= g								
FT	FT	/note= "OTHER = 2'-O-methyl nucleotide"								
FT	FT	21..27								
FT	FT	/*tag= c								
FT	FT	/mod_base= OTHER								
FT	FT	/note= "OTHER = 2'-O-methyl nucleotides"								
FT	FT	27								
FT	FT	/*tag= d								
FT	FT	/mod_base= c								
FT	FT	/note= "3',3'-inverted deoxyabasic moiety"								
XX	XX									
PN	WO200159102-A2.									
XX	PD	16-AUG-2001.								
XX	XX									
PF	08-FEB-2001;	2001WO-US04223.								
PR	08-FEB-2000;	2000US-0181360.								
PR	31-MAR-2000;	2000US-0193646.								
XX	(RIBO-) RIBOZYME PHARM INC.									
PA	(UYVA) UNIV YALE.									
XX	Breaker R, Beigelman L, Emilsson G;									
PI	WPI; 2001-536526/59.									
DR	XX									
XX	XX									
PT	XX	New nucleic acids with endonuclease activity, such as ribozymes and								
PT	XX	nucleozymes, for modulating gene expression in a plant, mammalian,								
PT	XX	bacterial or fungal cell -								
FS	Example 1; Page 71; 96pp; English.									
PS	XX									
CC	XX	The invention relates to nucleic acid molecules with endonuclease								
CC	CC	activity, which are particularly useful for cleavage of RNA or DNA.								
CC	CC	The nucleic acids are used in a pharmaceutical composition and are used								
CC	CC	to modulate expression of a gene in a plant, mammalian, bacterial or								
CC	CC	fungal cell. They are used to cleave a separate nucleic acid, preferably								
CC	CC	RNA. The nucleic acids are used to inhibit gene expression and/or cell								
CC	CC	proliferation, and can be used to treat a disease or condition. More								
CC	CC	than one nucleic acid can be independently targeted to the same or								
CC	CC	different sites in a cell. The nucleic acids may be used to study DNA.								
CC	CC	The modifications to the nucleic acids optimises their catalytic activity								
CC	CC	and can maintain or enhance their activity. They exhibit a high degree								

CC of specificity for RNA. The present sequence represents the coding
CC sequence of class V ribozyme #17 used in the method of the invention.

XX
SQ Sequence 27 BP; 11 A; 3 C; 9 G; 4 U; 0 other;
Query Match 100.0%; Score 15; DB 22; Length 27;
Best Local Similarity 100.0%; Pred. No. 74;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGUAUACGUGAAGAU 15
|||||
Db 7 AGUAUACGUGAAGAU 21

RESULT 13

AAS12306
ID AAS12306 standard; DNA; 27 BP.

XX
AC AAS12306;

XX
DT 21-NOV-2001 (first entry)

XX
DE DNA encoding class V ribozyme #18.

XX
KW Ribozyme; cytosolic; endonuclease; RNA cleavage; DNA cleavage;
KW gene therapy; plant; fungus; bacteria; mammal; ss.

XX
OS Synthetic.

XX
FH Key Location/Qualifiers
FT modified_base 1..6

FT /tag= a
FT /mod_base= OTHER
FT /note= "OTHER = 2'-O-methyl nucleotides"
FT modified_base 14

FT /tag= b
FT /mod_base= g
FT /note= "OTHER = 2'-O-methyl nucleotide"
FT modified_base 21..27

FT /tag= c
FT /mod_base= OTHER
FT /note= "OTHER = 2'-O-methyl nucleotides"
FT modified_base 27

FT /tag= d
FT /mod_base= c
FT /note= "3',3'-inverted deoxyabasic moiety"

XX
PN WO200159102-A2.

XX
PD 16-AUG-2001.

XX
PF 08-FEB-2001; 2001WO-US04223.

XX
PR 08-FEB-2000; 2000US-0181360.

XX
PR 31-MAR-2000; 2000US-0193646.

XX
PA (RIBO-) RIBOZYME PHARM INC.
PA (UYVA) UNIV YALE.

XX
PI Breaker R, Beigelman L, Emilsson G;

XX
WPI; 2001-536526/59.

XX
PT New nucleic acids with endonuclease activity, such as ribozymes and
PT nucleozymes, for modulating gene expression in a plant, mammalian,
PT bacterial or fungal cell

XX
PS Example 1; Page 71; 96pp; English.

XX
CC The invention relates to nucleic acid molecules with endonuclease
CC activity, which are particularly useful for cleavage of RNA or DNA.
CC The nucleic acids are used in a pharmaceutical composition and are used
CC to modulate expression of a gene in a plant, mammalian, bacterial or

CC fungal cell. They are used to cleave a separate nucleic acid, preferably
CC RNA. The nucleic acids are used to inhibit gene expression and/or cell
CC proliferation, and can be used to treat a disease or condition. More
CC than one nucleic acid can be independently targeted to the same or
CC different sites in a cell. The nucleic acids may be used to study DNA.
CC The modifications to the nucleic acids optimises their catalytic activity
CC and can maintain or enhance their activity. They exhibit a high degree
CC of specificity for RNA. The present sequence represents the coding
CC sequence of class V ribozyme #18 used in the method of the invention.

XX
SQ Sequence 27 BP; 11 A; 3 C; 9 G; 4 U; 0 other;

Query Match 100.0%; Score 15; DB 22; Length 27;

Best Local Similarity 100.0%; Pred. No. 74;

Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGUAUACGUGAAGAU 15

|||||

Db 7 AGUAUACGUGAAGAU 21

RESULT 14

AAS12307

ID AAS12307 standard; DNA; 27 BP.

XX
AC AAS12307;

XX
DT 21-NOV-2001 (first entry)

XX
DE DNA encoding class V ribozyme #19.

XX
KW Ribozyme; cytosolic; endonuclease; RNA cleavage; DNA cleavage;
KW gene therapy; plant; fungus; bacteria; mammal; ss.

XX
OS Synthetic.

XX
FH Key Location/Qualifiers
FT modified_base 1..6

FT /tag= a
FT /mod_base= OTHER
FT /note= "OTHER = 2'-O-methyl nucleotides"
FT modified_base 13

FT /tag= b
FT /mod_base= c
FT /note= "OTHER = 2'-O-methyl nucleotide"
FT modified_base 21..27

FT /tag= c
FT /mod_base= OTHER
FT /note= "OTHER = 2'-O-methyl nucleotides"
FT modified_base 27

FT /tag= d
FT /mod_base= c
FT /note= "3',3'-inverted deoxyabasic moiety"

XX
PN WO200159102-A2.

XX
PD 16-AUG-2001.

XX
PF 08-FEB-2001; 2001WO-US04223.

XX
PR 08-FEB-2000; 2000US-0181360.

XX
PR 31-MAR-2000; 2000US-0193646.

XX
PA (RIBO-) RIBOZYME PHARM INC.

XX
PA (UYVA) UNIV YALE.

XX
PI Breaker R, Beigelman L, Emilsson G;

XX
WPI; 2001-536526/59.

XX
PT New nucleic acids with endonuclease activity, such as ribozymes and
PT nucleozymes, for modulating gene expression in a plant, mammalian,
PT bacterial or fungal cell

XX PS Example 1; Page 71; 96pp; English.
XX CC The invention relates to nucleic acid molecules with endonuclease
CC activity, which are particularly useful for cleavage of RNA or DNA.
CC The nucleic acids are used in a pharmaceutical composition and are used
CC to modulate expression of a gene in a plant, mammalian, bacterial or
CC fungal cell. They are used to cleave a separate nucleic acid, preferably
CC RNA. The nucleic acids are used to inhibit gene expression and/or cell
CC proliferation, and can be used to treat a disease or condition. More
CC than one nucleic acid can be independently targeted to the same or
CC different sites in a cell. The nucleic acids may be used to study DNA.
CC The modifications to the nucleic acids optimises their catalytic activity
CC and can maintain or enhance their activity. They exhibit a high degree
CC of specificity for RNA. The present sequence represents the coding
CC sequence of class V ribozyme #19 used in the method of the invention.
XX SQ Sequence 27 BP; 11 A; 3 C; 9 G; 4 U; 0 other;
Query Match 100.0%; Score 15; DB 22; Length 27;
Best Local Similarity 100.0%; Pred. No. 74;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 AGAUAACGUGAAGAU 15
Db 7 AGAUAACGUGAAGAU 21
RESULT 15
AAS12308
ID AAS12308 standard; DNA; 27 BP.
XX AC AAS12308;
XX DT 21-NOV-2001 (first entry)
XX DE DNA encoding class V ribozyme #20.
XX KW Ribozyme; cytosstatic; endonuclease; RNA cleavage; DNA cleavage;
XX KW gene therapy; plant; fungus; bacteria; mammal; ss.
XX OS Synthetic.
XX FH Key Location/Qualifiers
FT modified_base 1..6 /*tag= a
FT /*mod_base= OTHER
FT /*note= "OTHER = 2'-O-methyl nucleotides"
FT modified_base 21..27 /*tag= b
FT /*mod_base= OTHER
FT /*note= "OTHER = 2'-O-methyl nucleotides"
FT modified_base 27 /*tag= c
FT /*mod_base= c
FT /*note= "3',3'-inverted deoxybasic moiety"
XX PN WO200159102-A2.
XX PD 16-AUG-2001.
XX PF 08-FEB-2001; 2001WO-US04223.
XX PR 08-FEB-2000; 2000US-0181360.
XX PR 31-MAR-2000; 2000US-0193646.
XX PA (RIBO-) RIBOZYME PHARM INC.
XX PA (UYA) UNIV YALE.
XX PI Breaker R, Beigelman L, Emilsson G;
XX DR WPI; 2001-536526/59.
XX

PT New nucleic acids with endonuclease activity, such as ribozymes and
PT nucleozymes, for modulating gene expression in a plant, mammalian,
PT bacterial or fungal cell
XX Example 1; Page 71; 96pp; English.
XX CC The invention relates to nucleic acid molecules with endonuclease
CC activity, which are particularly useful for cleavage of RNA or DNA.
CC The nucleic acids are used in a pharmaceutical composition and are used
CC to modulate expression of a gene in a plant, mammalian, bacterial or
CC fungal cell. They are used to cleave a separate nucleic acid, preferably
CC RNA. The nucleic acids are used to inhibit gene expression and/or cell
CC proliferation, and can be used to treat a disease or condition. More
CC than one nucleic acid can be independently targeted to the same or
CC different sites in a cell. The nucleic acids may be used to study DNA.
CC The modifications to the nucleic acids optimises their catalytic activity
CC and can maintain or enhance their activity. They exhibit a high degree
CC of specificity for RNA. The present sequence represents the coding
CC sequence of class V ribozyme #20 used in the method of the invention.
XX SQ Sequence 27 BP; 11 A; 3 C; 9 G; 4 U; 0 other;
Query Match 100.0%; Score 15; DB 22; Length 27;
Best Local Similarity 100.0%; Pred. No. 74;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 AGAUAACGUGAAGAU 15
Db 7 AGAUAACGUGAAGAU 21
Search completed: July 6, 2003, 14:32:52
Job time : 160.909 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2003 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: July 6, 2003, 14:26:51 : Search time 1007.73 Seconds
(without alignments)
241.069 Million cell updates/sec

Title: US-09-780-929-97

Perfect score: 15

Sequence: 1 agaaacgugaagau 15

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 16154066 seqs, 8097743376 residues

Total number of hits satisfying chosen parameters: 146654

Minimum DB seq length: 0

Maximum DB seq length: 60

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

EST:*

1: em_estba:.*
2: em_esthum:.*
3: em_estin:.*
4: em_estnu:.*
5: em_estov:.*
6: em_estpl:.*
7: em_estro:.*
8: em_htc:.*
9: gb_estl:.*
10: gb_est2:.*
11: gb_htc:.*
12: gb_est3:.*
13: gb_est4:.*
14: gb_est5:.*
15: em_estfun:.*
16: em_eston:.*
17: gb_gss:.*
18: em_gss_hum:.*
19: em_gss_inv:.*
20: em_gss_pln:.*
21: em_gss_vrt:.*
22: em_gss_fun:.*
23: em_gss_mam:.*
24: em_gss_mus:.*
25: em_gss_other:.*
26: em_gss_pro:.*
27: em_gss_rod:.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Match	Length	DB ID	Description
C 1	11.8	78.7	29	17	AZ994980
2	11.8	78.7	59	14	BQ625912
3	10.8	72.0	22	17	TA248D01P
4	10.8	72.0	28	17	AZ345853
5	10.8	72.0	37	9	AA947987
6	10.8	72.0	37	17	AL760544
					AL760544 Arabidops

C	7	10.8	72.0	40	9	AI039249
	8	10.8	72.0	46	17	AZ482955
	9	10.8	72.0	48	17	AZ656367
	10	10.8	72.0	48	17	AZ772291
C	11	10.8	72.0	50	9	AU103231
	12	10.8	72.0	51	10	AV833384
	13	10.8	72.0	52	9	AA823664
C	14	10.8	72.0	52	12	BF631905
	15	10.8	72.0	53	9	AU257573
C	16	10.8	72.0	54	13	BM069548
C	17	10.8	72.0	54	17	AF149434
	18	10.8	72.0	58	9	AA780323
	19	10.8	72.0	59	13	BJ053305
	20	10.4	69.3	42	13	BJ044345
	21	10.4	69.3	44	14	D18694
	22	10.4	69.3	48	14	D18688
	23	10.4	69.3	51	10	AW100987
	24	10.4	69.3	53	17	AL764237
C	25	10.4	69.3	55	10	AW424064
	26	10.4	69.3	56	12	BG731546
C	27	10.4	69.3	56	17	AZ784261
	28	10.4	69.3	57	9	AU256712
	29	10.4	69.3	58	9	AA265174
C	30	10.4	69.3	60	13	BI493320
C	31	10.4	69.3	60	17	AL752372
	32	10.2	68.0	19	17	AZ759898
C	33	10.2	68.0	24	17	TA200H11P
C	34	10.2	68.0	33	17	AL760055
C	35	10.2	68.0	37	17	AZ760010
C	36	10.2	68.0	40	9	AI609205
C	37	10.2	68.0	40	9	AI638565
	38	10.2	68.0	40	14	N54453
	39	10.2	68.0	42	17	AZ388234
C	40	10.2	68.0	42	17	AZ449920
C	41	10.2	68.0	44	12	BE738334
	42	10.2	68.0	45	17	BH638406
	43	10.2	68.0	46	9	AI077563
C	44	10.2	68.0	46	17	AZ321153
C	45	10.2	68.0	48	17	BH863621

ALIGNMENTS

RESULT 1
AZ994980/c
LOCUS
DEFINITION 2M0280G07R Mouse 10kb plasmid UUC2M library Mus musculus genomic clone UUC2M0280G07 R, DNA sequence.
ACCESSION AZ994980
VERSION AZ994980.1 GI:13866207
KEYWORDS GSS.
SOURCE house mouse.
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE 1 (bases 1 to 29)
AUTHORS Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamil, C., Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T., Reilly, M., Rose, M., Rose, R., Stokes, R., Tingey, A., von Niederhausern, A., and Wright, D., Weiss, R.
TITLE Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts
JOURNAL Unpublished (2000)
COMMENT Contact: Robert B. Weiss
University of Utah Genome Center
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT 84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00

AZ994980 29 bp DNA linear GSS 27-APR-2001
2M0280G07R Mouse 10kb plasmid UUC2M library Mus musculus genomic clone UUC2M0280G07 R, DNA sequence.

AZ994980 GI:13866207

GSS.

house mouse.

Mus musculus

Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

1 (bases 1 to 29)

Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamil, C.,

Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T., Reilly

, M., Rose, M., Rose, R., Stokes, R., Tingey, A., von Niederhausern, A.

, and Wright, D., Weiss, R.

Mouse whole genome scaffolding with paired end reads from 10kb

plasmid inserts

Unpublished (2000)

Contact: Robert B. Weiss

University of Utah Genome Center

Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT

84112, USA

Tel: 801 585 5606

Fax: 801 585 7177

Email: ddunn@genetics.utah.edu

Insert Length: 10000 Std Error: 0.00

Plate: 0280 row: G column: 07
Seq primer: CACACAGGAACACCTATGACC

Class: plasmid ends
High quality sequence stop: 29.

FEATURES

Location/Qualifiers
1. .29
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUGC2M0280G07"
/clone_lib="Mouse 10kb plasmid UUGC2M library"
/sex="Female"
/lab_host="E. coli strain XL10-Gold, T1-resistant, F-"
/note="Vector: pWD42nv; Purified genomic DNA from M. Laboratory Mouse DNA Resource

musculus C57BL/6J (female) was obtained from the Jackson
http://www.jax.org/resources/documents/dnares/). The DNA
was hydrodynamically sheared by repeated passage through a
0.005 inch orifice at constant velocity. The sheared DNA
was blunt end-repaired with T4 DNA polymerase and T4
polynucleotide kinase. Adaptor oligonucleotides were
ligated to the blunt ends in high molar excess. The
adaptor DNA was purified and size-selected for a 9.5 to
10.5 kb range using preparative agarose gel
electrophoresis. Vector DNA was prepared from a derivative
of pWD42 (gi4732114|gb|AF129072.1), a copy-number
inducible derivative of plasmid RL. The vector was ligated
with adaptors complementary to the insert adaptors and
purified. The sheared, adaptor-muscle DNA was annealed to
adaptor vector DNA, and transformed into
chemically-competent E. coli XL10-Gold (Stratagene) cells
and selected for ampicillin resistance."

BASE COUNT 5 a 5 c 5 g 14 t

Query Match 78.7%; Score 11.8; DB 17; Length 29;
Best Local Similarity 66.7%; Pred. No. 1.2e+04;
Matches 10; Conservative 3; Mismatches 2; Indels 0; Gaps 0;

Qy 1 AGAUACGUGAAGAU 15
||||| 1:|||||
Db 23 AGATAGAGTGAAGAT 9

RESULT 2

BQ625912
LOCUS
DEFINITION
ph86f02.v1 Ostertagia ostertagi L3 SLL TOPO v2 Ostertagia ostertagi
CDNA 5', mRNA sequence.

ACCESSION BQ625912
VERSION BQ625912.1 GI:21653090
KEYWORDS EST.

SOURCE

ORGANISM
Ostertagia ostertagi.
Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida; Strongylida;
Trichostrongylidae; Haemonchidae; Ostertagiinae; Ostertagia.

1 (bases 1 to 59)

McCartner, J., Clifton, S., Chiapelli, B., Pape, D., Martin, J., Wylie, T.,
Dante, M., Marra, M., Hillier, L., Kucaba, T., Theising, B., Bowers, Y.,
Gibbons, M., Ritter, E., Bennett, J., Franklin, C., Tsagarisvili, R.,
Ronko, I., Kennedy, S., Maguire, L., Beck, C., Underwood, K., Steptoe
M., Allen, M., Person, B., Swaller, T., Harvey, N., Schurk, R., Kohn, S.,
Shin, T., Jackson, Y., Cardenas, M., McCann, R., Waterston, R. and
Wilson, R.

The Washington Univ. Nematode EST Project, 1999
Unpublished (1999)

Contact: McCartner JP

The Washington Univ. Nematode EST Project, 1999
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA

Tel: 314 286 1800

Fax: 314 286 1810

Email: est@watson.wustl.edu

FEATURES

source

Seq primer: SLL primer.

The vector to vector length is 60

Location/Qualifiers

1. .59

/organism="Ostertagia ostertagi"

/db_xref="taxon:6317"

/clone_lib="Ostertagia ostertagi L3 SLL TOPO v2"

/dev_stage="third stage exsheathed larvae"

/lab_host="DH10B"

/note="Vector: pCRII-TOPO (Invitrogen); Site_1: EcoRI;
Site_2: EcoRI; The library was constructed by Claire
Murphy and Dr. James McCartner at Washington University,
St. Louis. Oligo(dT)-SLL PCR based library. Ostertagia
ostertagi L3 cDNA PCR products of size >400 nucleotides
containing SLL on the 5' end and oligo(dT) on the 3' end
were non-directionally cloned into pCRII-TOPO(Invitrogen)
following the TOPO TA cloning protocol. Nematodes were
provided by Dr. Louis Gasbarre of the USDA, Beltsville,
MD (lgasbarre@nri.barc.usda.gov). Third stage exsheathed
larvae were collected from 14 day fecal-sphagnum moss
cultures of Ostertagia eggs. The larvae were recovered by
overnight passage on a Baermann apparatus, and then
cleaned by passage through a 20 micron nylon mesh. The
larvae were then subjected to a treatment with 1.25%
chlorox to induce excystation. The larvae were washed with
5 changes of PBS and then pelleted and snap frozen in
liquid nitrogen."

BASE COUNT 22 a 16 c 12 g 9 t

Query Match 78.7%; Score 11.8; DB 14; Length 59;

Best Local Similarity 73.3%; Pred. No. 1.7e+04;

Matches 11; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

Qy 1 AGAUACGUGAAGAU 15
||||| 1:|||||
Db 7 AGAAAAGGTGAAGAT 21

RESULT 3

TA248D01P/c

LOCUS

DEFINITION

T. brucei sheared genomic DNA clone 248d01, forward sequence,

genomic survey sequence.

ACCESSION AL483187

VERSION AL483187.1 GI:11848863

KEYWORDS GSS.

SOURCE Trypanosoma brucei.

ORGANISM

Trypanosoma brucei

Eukaryota; Euglenozoa; Kinetoplastida; Trypanosomatidae;

Trypanosoma.

1 (bases 1 to 22)

Hall, N., Bowman, S., Lennard, N.J., Doggett, J., Atkin, R.,
Chillingworth, C., Ormond, D., Harris, B., El-Sayed, N., Hou, L.,
Melville, S.E., Rajandream, M.A. and Barrell, B.G.

Direct Submission

Submitted (10-DEC-2000)

Project, Sanger Centre, The Wellcome Trust Genome Campus, Hinxton,

Cambridge CB10 1SA, E-mail: barrellesanger.ac.uk and

nhl@sanger.ac.uk

COMMENT

Constructed at the Institute for Genomic Research (TIGR),

Rockville, MD. Genomic DNA isolated from a cloned population of

Trypanosoma brucei (TREU927/4 GUTAT 10.1) was mechanically sheared

to give a tight size distribution (

4 kb). The v + i method used for the library construction is

described in detail in Smith, H. and Venter, J.C. (Making small

insert libraries for whole genome shotgun sequencing projects. In

Genome Sequencing: A Practical Approach, eds. M. Vaudin and B.

The library was constructed by Claire Murphy and Dr. James McCartner
at Washington University, St. Louis. Nematodes were provided by Dr.
Louis Gasbarre of the USDA, Beltsville, MD
(lgasbarre@nri.barc.usda.gov).
Putative full length read

Barrell, Oxford University Press, 1999).

Email: nelsayed@tigr.org
Details of T. brucei sequencing at the Sanger Centre are available
at http://www.sanger.ac.uk/Projects/T_brucei/.

FEATURES

Location/Qualifiers
1..22

/organism="Trypanosoma brucei"
/strain="TREU927"
/db_xref="taxon:5691"
/clone="248d01"

BASE COUNT 5 a 6 c 2 g 9 t

ORIGIN

Query Match 72.0%; Score 10.8; DB 17; Length 22;
Best Local Similarity 64.3%; Pred. No. 3.8e+04; Indels 0; Gaps 0;
Matches 9; Conservative 3; Mismatches 2; Indels 0; Gaps 0;

QY 2 GAUAACGUGAAGAU 15

DB 19 GATACATGAAAT 6

RESULT 4

AZ345853

LOCUS

DEFINITION 1M0080F14R Mouse 10kb plasmid UUC1M library Mus musculus genomic
clone UUC1M0080F14 R, DNA sequence.

ACCESSION AZ345853

VERSION AZ345853.1 GI:10425090

KEYWORDS

GSS.

SOURCE house mouse.

ORGANISM Mus musculus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

REFERENCE

AUTHORS 1 (bases 1 to 28)
Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C.,
Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly
M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A.
and Wright,D., Weiss,R.

TITLE Mouse whole genome scaffolding with paired end reads from 10kb
plasmid inserts

JOURNAL Unpublished (2000)

COMMENT Contact: Robert B. Weiss

University of Utah Genome Center

Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT

84112, USA

Tel: 801 585 5606

Fax: 801 585 7177

Email: ddunn@genetics.utah.edu

Insert Length: 10000 Std Error: 0.00

Plate: 0080 row: F column: 14

Seq primer: CACACAGGAACAGCTATGACC

Class: plasmid ends

High quality sequence stop: 28.

Location/Qualifiers

FEATURES

source

1..28

/organism="Mus musculus"

/strain="C57BL/6J"

/db_xref="taxon:10090"

/clone="UUC1M0080F14"

/clone_lib="Mouse 10kb plasmid UUC1M library"

/sex="Male"

/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"

/note="Vector: PWD42nv; Purified genomic DNA from M.

musculus C57BL/6J (male) was obtained from the Jackson

Laboratory Mouse DNA Resource

(<http://www.jax.org/resources/documents/dnares/>). The DNA

was hydrodynamically sheared by repeated passage through a

0.005 inch orifice at constant velocity. The sheared DNA

was blunt end-repaired with T4 DNA polymerase and T4

polynucleotide kinase. Adaptor oligonucleotides were

ligated to the blunt ends in high molar excess. The

adapted DNA was purified and size-selected for a 9.5 to
10.5 kb range using preparative agarose gel
electrophoresis. Vector DNA was prepared from a derivative
of pWD42 (g114732114|g1b1AF129072.1), a copy-number
inducible derivative of plasmid RL. The vector was ligated
with adaptors complementary to the insert adaptors and
purified. The sheared, adapted mouse DNA was annealed to
adapted vector DNA, and transformed into
chemically-competent E. coli XL10-Gold (Stratagene) cells
and selected for ampicillin resistance.

BASE COUNT 16 a 3 c 4 g 5 t

ORIGIN

Query Match 72.0%; Score 10.8; DB 17; Length 28;
Best Local Similarity 71.4%; Pred. No. 4.3e+04; Indels 0; Gaps 0;
Matches 10; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 1 AGAUAACGUGAAGA 14

DB 9 AAATAACGTGAAA 22

RESULT 5

AA947987

LOCUS

DEFINITION AA947987 37 bp mRNA linear EST 04-MAY-1998
Oq58e02.s1 NCL_CGAP_Kid6 Homo sapiens cDNA clone IMAGE:1590554 3'
similar to TR:Q33559 Q33559 NH2 TERMINUS UNCERTAIN ;, mRNA
sequence.

ACCESSION AA947987

VERSION AA947987.1 GI:3109240

KEYWORDS

EST..

SOURCE human.

ORGANISM

Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE

1 (bases 1 to 37)

AUTHORS

NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.

TITLE

National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index

JOURNAL Unpublished (1997)

COMMENT Contact: Robert Strausberg, Ph.D.

Email: cgapbs-r@mail.nih.gov

Tissue Procurement: L. Jeffrey Medeiros, M.D., Michael R.

Emmert-Buck, M.D., Ph.D.

cDNA Library Preparation: Stratagene, Inc.

cDNA Library Arrayed by: Greg Lennon, Ph.D.

DNA Sequencing by: Washington University Genome Sequencing Center

Clone distribution: NCI-CGAP clone distribution information can be

found through the I.M.A.G.E. Consortium/LLNL at:

www-bio.llnl.gov/bbrp/image/image.html

Trace considered overall poor quality

Seq primer: -40ml3 fwd. ET from Amersham

High quality sequence stop: 1.

Location/Qualifiers

1..37

/organism="Homo sapiens"

/db_xref="taxon:9606"

/clone="IMAGE:1590554"

/clone_lib="NCI-CGAP_Kid6"

/sex="mixed"

/tissue_type="kidney tumor"

/lab_host="SOLR (kanamycin resistant)"

/note="Organ: Kidney; Vector: Bluescript SK-; Site:1;

EcoRI; Site:2; XhoI: Cloned unidirectionally. Primer:

Oligo dr. Pooled kidney tumors. 5' adaptor sequence: 5'

GAATTCGGCAGAG 3' 3' adaptor sequence: 5'

CATCGAGTTTTTTTTTTT 3' Average insert size: 1.0 kb.

BASE COUNT 26 a 2 c 6 g 3 t

ORIGIN

Query Match 72.0%; Score 10.8; DB 9; Length 37;

Best Local Similarity 78.6%; Pred. No. 4.9e+04;

```

Matches 11; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

QY 1 AGAUAAACGUGAAGA 14
Db 17 AGATAATGAGAAGA 30

RESULT 6
AL760544
LOCUS
DEFINITION
Arabidopsis thaliana T-DNA flanking sequence GK-199B11-014765,
genomic survey sequence.
ACCESSION
AL760544
VERSION
AL760544.1 GI:21499415
KEYWORDS
GSS.
SOURCE
thale cress.
ORGANISM
Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
Rosidae; eurosids II; Brassicales; Brassicaceae; Arabidopsids.
REFERENCE
1 Strizhov,N., Li,Y., Rosso,M., Viehoever,P., Dekker,K., Saedler,H.
and Weisshaar,B.
TITLE
A pipeline for automated high-throughput generation of ESTs
(flanking sequence tags) from Arabidopsis thaliana T-DNA
transformed lines
JOURNAL
Unpublished
REFERENCE
2 Rosso,M., Strizhov,N., Li,Y., Reiss,B., Dekker,K. and Weisshaar,B.
A new Arabidopsis thaliana T-DNA mutagenised population (GABI-Kat)
for flanking sequence tag based reverse genetics
JOURNAL
Unpublished
REFERENCE
3 (bases 1 to 37)
Li,Y., Strizhov,N., Rosso,M. and Weisshaar,B.
Direct Submission
Submitted (17-JUN-2002) Weisshaar B., Max-Planck-Institut fuer
Zuechtungsforschung, Carl-von-Linne-Weg 10, Koeln, 50829, Germany
This sequence is recovered from the left border of the T-DNA. It
indicates an insertion close to or within gene Atig77800. The
sequences are generated at the MPI for Plant Breeding Research in
the context of the GABI-Kat project. GABI-Kat is part of the German
Plant Genomics program designated 'GABI'. Information on line
availability can be found at:
http://www.mpiz-koeln.mpg.de/GABI-Kat/.
FEATURES
Location/Qualifiers
1..37
/organism="Arabidopsis thaliana"
/strain="Columbia 0"
/db_xref="taxon:3702"
/clone="GK-199B11-014765"
/note="PCR was performed on DNA from Arabidopsis thaliana
plants (T1) which were transformed with the T-DNA from
vector PAC161. The lines contain one or more T-DNA
insertions. The DNA fragment(s) resulting from the PCR
were directly sequenced to determine the genomic sequence
flanking the insertion. Sequences displaying significant
similarity to the A. thaliana nuclear genome sequence were
processed for submission. T-DNA derived sequences were
removed"
BASE COUNT 19 a 4 c 6 g 8 t
ORIGIN
Query Match 72.0%; Score 10.8; DB 17; Length 37;
Best Local Similarity 78.6%; Pred. No. 4.9e+04;
Matches 11; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

QY 1 AGAUAAACGUGAAGA 14
Db 21 AGAAAACATGAAGA 34

RESULT 7

```

AI039249/c

LOCUS

DEFINITION

AI039249 40 bp mRNA linear EST 30-JUN-1998
 ox33a08.s1 Soares total_fetus.Nb2Hf8_9w Homo sapiens cDNA clone
 IMAGE:1658102 3' similar to TR:000554 000554 P21-ARC..[1] ;, mRNA
 sequence.

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

FEATURES

Source

1..40

/organism="Homo sapiens"

/db_xref="taxon:9606"

/clone="IMAGE:1658102"

/dev_stage="8-9 weeks"

/lab_host="DH10B"

/note="Vector: pT7T3D-Pac (Pharmacia) with a modified

polylinker; Site_1: Not I; Site_2: Eco RI; 1st strand

was prepared from mRNA obtained from pooled 8-9 week

(total) fetus material with a Not I - oligo(dT) primer [5'

TGTTACCAATCTGAAGTGGAGCGCGCTTAATTTTTTTTTTTT 3'].

Double-stranded cDNA was ligated to Eco RI adaptors

(Pharmacia), digested with Not I and cloned into the Not I

and Eco RI sites of the modified pT7T3 vector. Library

went through one round of normalization, and was

constructed by Bento Soares and M. Fatima Bonaldo. "

BASE COUNT 3 a 11 c 10 g 16 t

ORIGIN

Query Match 72.0%; Score 10.8; DB 9; Length 40;

Best Local Similarity 71.4%; Pred. No. 5.1e+04;

Matches 10; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 1 AGAUAAACGUGAAGA 14

Db 17 AGATGACGTGATGA 4

RESULT 8

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

1 (bases 1 to 46)

Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamill,C.,

Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly

M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A.

and Wright,D., Weiss,R.

Mouse whole genome scaffolding with paired end reads from 10kb

plasmid inserts

plasmid inserts

AZ482955

IM0308M13F Mouse 10kb plasmid UUGC1M library Mus musculus genomic

clone UUGC1M0308M13 F, DNA sequence.

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

1 (bases 1 to 46)

Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamill,C.,

Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly

M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A.

and Wright,D., Weiss,R.

Mouse whole genome scaffolding with paired end reads from 10kb

plasmid inserts

JOURNAL
COMMENT

Unpublished (2000)
Contact: Robert B. Weiss
University of Utah
Genome Center
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLIC, UT
84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0308 row: M column: 13
Seq primer: CGTGTAAACAGCAGCGCCAGT
Class: plasmid ends
High quality sequence stop: 46.
Location/Qualifiers
1. 46

FEATURES
SOURCE

/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUGC1M0308M13"
/clone_lib="Mouse 10kb plasmid UUGC1M library"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, Tl-resistant, F-"
/note="Vector: PWD42nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource (http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adapted DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of pWD42 (gil147321141gb|AF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adapted mouse DNA was annealed to adapted vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT
ORIGIN

26 a 6 c 8 g 6 t
Query Match 72.0%; Score 10.8; DB 17; Length 46;
Best Local Similarity 78.6%; Pred. No. 5.4e+04;
Matches 11; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

QY 1 AGAUAACGUGAAGA 14
| | | | | | | | | | | | | | | |
DB 19 AGAAACGTGAAAA 32

RESULT 9
AZ656367

LOCUS 48 bp DNA linear GSS 14-DEC-2000
DEFINITION IM0531D23R Mouse 10kb plasmid UUGC1M library Mus musculus genomic clone UUGC1M0531D23 R, DNA sequence.

ACCESSION
VERSION
KEYWORDS
SOURCE

AZ656367
GI:11793513

ORGANISM
house mouse.
Mus musculus

Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus. 1 (bases 1 to 48)

REFERENCE
AUTHORS

Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C., Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A. and Wright,D., Weiss,R.

Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts

JOURNAL
COMMENT

Unpublished (2000)
Contact: Robert B. Weiss
University of Utah
Genome Center
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLIC, UT
84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0531 row: D column: 23
Seq primer: CACACAGGAACAGCTATGACC
Class: plasmid ends
High quality sequence stop: 48.
Location/Qualifiers
1. 48

FEATURES
SOURCE

/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUGC1M0531D23"
/clone_lib="Mouse 10kb plasmid UUGC1M library"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, Tl-resistant, F-"
/note="Vector: PWD42nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource (http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adapted DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of pWD42 (gil147321141gb|AF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adapted mouse DNA was annealed to adapted vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

26 a 8 c 6 g 8 t

BASE COUNT
ORIGIN

Query Match 72.0%; Score 10.8; DB 17; Length 48;
Best Local Similarity 71.4%; Pred. No. 5.6e+04;
Matches 10; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 1 AGAUAACGUGAAGA 14
| | | | | | | | | | | | | | | |
DB 20 AAATAACGTGAAAA 33

RESULT 10
AZ772291

LOCUS 48 bp DNA linear GSS 16-FEB-2001
DEFINITION IM0583L06F Mouse 10kb plasmid UUGC1M library Mus musculus genomic clone UUGC1M0583L06 F, DNA sequence.

ACCESSION
VERSION
KEYWORDS
SOURCE

AZ772291
GI:12895445

ORGANISM
house mouse.
Mus musculus

Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus. 1 (bases 1 to 48)

REFERENCE
AUTHORS

Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C., Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A. and Wright,D., Weiss,R.

Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts

JOURNAL
COMMENT

Unpublished (2000)
Contact: Robert B. Weiss
University of Utah Genome Center
University of Utah
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT
84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: dunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0583 row: L column: 06
Seq primer: CGTTGTAACACGACGCCAGT
Class: plasmid ends
High quality sequence stop: 48.

FEATURES

source

1. .48
Location/Qualifiers
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUGC1M0583106"
/clone_lib="Mouse 10kb plasmid UUGCLM library"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
/note="Vector: pWD42nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource (http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of pWD42 (g147321141gb)AF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT 23 a 8 c 7 g 10 t
ORIGIN

Query Match 72.0%; Score 10.8; DB 17; Length 48;
Best Local Similarity 71.4%; Pred. No. 5.6e+04;
Matches 10; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 1 AGUAACGUGAGA 14
Db 33 AAATAACGTGAAAA 46

RESULT 11

AU103231/c
LOCUS AU103231 Sugano Homo sapiens cDNA library EST 30-AUG-2001
DEFINITION KAT07952, mRNA sequence.
ACCESSION AU103231
VERSION AU103231.1 GI:13552752
KEYWORDS EST.
SOURCE human.

ORGANISM

Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE

1 (bases 1 to 50)
Suzuki,Y., Taira,H., Tsunoda,T., Mizushima-Sugano,J., Sese,J., Hata
H., Ota,T., Isogai,T., Tanaka,T., Morishita,S., Okubo,K., Sakaki
Y., Nakamura,Y., Suyama,A. and Sugano,S.

TITLE

Diverse transcriptional initiation revealed by fine, large-scale
mapping of mRNA start sites

JOURNAL

EMBO Rep. 2 (5), 388-393 (2001)

MEDLINE
COMMENT

21270072
Contact: Yutaka Suzuki
Department of Virology
Institute of Medical Science, University of Tokyo
4-6-1, Shirokanedai, Minatoku, Tokyo 108-8639, Japan
Email: ysuzuki@ims.u-tokyo.ac.jp
Suzuki,Y., Yoshitomo-Nakagawa,K., Maruyama,K., Suyama,A. and Sugano
S. Construction and characterization of a full length-enriched and
a 5'-end-enriched cDNA library. Gene 200 (1-2), 149-156 (1997).

FEATURES

source

1. .50
Location/Qualifiers
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="KAT07952"
/clone_lib="Sugano Homo sapiens cDNA library"
/note="Differential display comparison of untreated and
dimethylfumarate treated U937 cells"
BASE COUNT 17 a 12 c 9 g 12 t
ORIGIN

Query Match 72.0%; Score 10.8; DB 9; Length 50;
Best Local Similarity 71.4%; Pred. No. 5.7e+04;
Matches 10; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 2 GAUAACGUGAAGAU 15
Db 46 GAGATGTGAAGAT 33

RESULT 12

AV833384
LOCUS AV833384 K. Sato unpublished cDNA library: Hordeum vulgare subsp.
DEFINITION AV833384 K. Sato unpublished cDNA library: Hordeum vulgare subsp.
ACCESSION AV833384.1 GI:14525473
VERSION AV833384
KEYWORDS EST.
SOURCE Hordeum vulgare subsp. vulgare.
ORGANISM Hordeum vulgare subsp. vulgare.
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Poideae
; Triticeae; Hordeum.

REFERENCE

1 (bases 1 to 51)
Sato,K.

AUTHORS Barley EST sequencing project in NIG and Okayama Univ
TITLE Unpublished (2001)
JOURNAL

COMMENT

Contact: Kazuhiro Sato
Research Institute for Bioresources
Okayama University, Barley Germplasm Center
Chuo 2-20-1, Kurashiki, Okayama 710-0046, Japan
Email: kassato@rib.okayama-u.ac.jp,
URL: http://www.rib.okayama-u.ac.jp/barley/
Sato,K., Saisho,D., Takeda,K., Shini,T. and Kohara,Y. Direct
submission;
database: http://www.shigen.nig.ac.jp/barley/Barley.html.

FEATURES

source

1. .51
Location/Qualifiers
/organism="Hordeum vulgare subsp. vulgare"
/cultivar="Haruna Nijo"
/db_xref="taxon:112509"
/clone="bags7123"
/clone_lib="K. Sato unpublished cDNA library: Hordeum
vulgare subsp. vulgare shoots germination"
/tissue_type="shoots"
/dev_stage="germination"
BASE COUNT 13 a 14 c 18 g 6 t
ORIGIN

Query Match 72.0%; Score 10.8; DB 10; Length 51;
Best Local Similarity 78.6%; Pred. No. 5.7e+04;

Matches 11; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

```

Qy      1 AGAUACGUGAAGA 14
      ||| ||||| |||
Db      36 AGAGACGCTGGAGA 49

RESULT 13
AA823664
LOCUS      52 bp      mRNA      linear      EST 17-FEB-1998
DEFINITION vt69d09.s1 Knowles Solter mouse 2 cell Mus musculus cDNA clone
IMAGE:1125905 5' similar to gb:J03161 SERUM RESPONSE FACTOR (HUMAN
);, mRNA sequence.
ACCESSION  AA823664
VERSION     AA823664.1 GI:2893532
KEYWORDS    EST.
SOURCE      Mus musculus
            house mouse.
ORGANISM    Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE   1 (bases 1 to 52)
AUTHORS     Maria.M., Hillier,L., Allen,M., Bowles,M., Dietrich,N., Dubuque,T.,
            Geisel,S., Kucaba,T., Lacy,M., Le,M., Martin,J., Morris,M.,
            Schellenberg,K., Steptoe,M., Tan,F., Underwood,K., Moore,B.,
            Theising,B., Wylie,T., Lennon,G., Soares,B., Wilson,R. and
            Waterston,R.
TITLE       The WashU-HHMI Mouse EST Project
JOURNAL     Unpublished (1996)
COMMENT     Contact: Marra M/Mouse EST Project
            WashU-HHMI Mouse EST Project
            Washington University School of MedicineP
            4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
            Tel: 314 286 1800
            Fax: 314 286 1810
            Email: mouseest@watson.wustl.edu
            This clone is available royalty-free through LLNL ; contact the
            IMAGE Consortium (info@image.llnl.gov) for further information.
            MGI:615241.

FEATURES             Location/Qualifiers
     source           1..52
     /organism="Mus musculus"
     /strain="C57BL/6J x DBA/2J F1"
     /db_xref="taxon:10090"
     /clone="IMAGE:1125905"
     /clone_lib="Knowles Solter mouse 2 cell"
     /tissue_type="embryo"
     /dev_stage="2-cell"
     /lab_host="DH10B"
     /note="Organ: embryo; Vector: pBluescribe (modified);
            Site 1: MluI; Site 2: SalI; Cloned unidirectionally from
            mRNA prepared from 13,500 2-cell stage embryos. Primer:
            SalI(drr): 5'-CGGTCGACCGTCGACCGTTTCTTTT-3'. cDNAs
            were cloned into the MluI/SalI sites of a modified
            pBluescribe vector using commercial linkers (NEB).
            Average insert size: 1.2 kb."
BASE COUNT      26 a      3 c      6 g      17 t
ORIGIN
      1
      2
      3
      4
      5
      6
      7
      8
      9
      10
      11
      12
      13
      14
      15
      16
      17
      18
      19
      20
      21
      22
      23
      24
      25
      26
      27
      28
      29
      30
      31
      32
      33
      34
      35
      36
      37
      38
      39
      40
      41
      42
      43
      44
      45
      46
      47
      48
      49
      50
      51
      52

Query Match      72.0%; Score 10.8; DB 9; Length 52;
Best Local Similarity 71.4%; Pred. NO. 5.8e+04;
Matches 10; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

Qy      1 AGAUACGUGAAGA 15
      ||| ||| ||| |||
Db      48 GATTAAGAGAGAT 35

RESULT 15
AA257573
LOCUS      53 bp      mRNA      linear      EST 25-APR-2002
DEFINITION AU257573 3'-directed mouse cDNA library Mus musculus cDNA clone
BED0010931 3', mRNA sequence.
ACCESSION  AU257573
VERSION     AU257573.1 GI:20322326
KEYWORDS    EST.
SOURCE      house mouse.
            Mus musculus
ORGANISM    Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE   1 (bases 1 to 53)
AUTHORS     Kato,K and Matoba,R.
TITLE       Generation of expressed sequence tags from mouse brain
JOURNAL     Unpublished (2002)
COMMENT     Contact: Kikuya Kato
            Graduate School of Biological Sciences
            Nara Institute of Science and Technology
            8916-5 Takayama, Ikoma, Nara 630-0101, Japan
            Tel: 81-743-72-5581
            Fax: 81-743-72-5589
            Email: kkatob@ns.aist-nara.ac.jp,
            URL:http://love2.aist-nara.ac.jp/BED/index.html.
            Location/Qualifiers
     source           1..53
     /organism="Mus musculus"
     /db_xref="taxon:10090"

```

SOURCE
ORGANISM

barrel medic.
Medicago truncatula
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
Rosidae; eurosids 1; Fabales; Fabaceae; Papilionoideae; Trifolieae;
Medicago.

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

1 (bases 1 to 52)

Torres-Jerez, I., Scott, A.D., Harris, A.R., Gonzales, R.A., Bell, C.J.,

Flores, H.R., Inman, J.T., Weller, J.W. and May, G.D.

Expressed Sequence Tags from the Samuel Roberts Noble Foundation

Medicago truncatula drought library

Unpublished (2000)

Contact: May GD

Plant Biology Division

The Samuel Roberts Noble Foundation

2510 Sam Noble Parkway, Ardmore, OK 73402, USA

Tel: 580 221 7391

Fax: 580 221 7380

Email: gdmay@noble.org

Insert length: 52 Std Error: 0.00

Plate: 016 Row: H Column: 07

Seq primer: TCACACGAGAAACAGCTATGAC.

Location/Qualifiers

1..52

/organism="Medicago truncatula"

/db_xref="taxon:3880"

/clone="NF016H07DT"

/clone_lib="Drought"

/tissue_type="Plantlets"

/dev_stage="Pooled timepoints"

/note="Vector: Lambda Zap; Contains a mixture of entire

plantlets harvested in a series of days-post-watering

timepoints."

11 a 15 c 5 g 21 t

BASE COUNT

ORIGIN

Query Match

Best Local Similarity

Matches 10; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

Qy

Db

RESULT 15

AA257573

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

1 (bases 1 to 53)

Kato, K and Matoba, R.

Generation of expressed sequence tags from mouse brain

Unpublished (2002)

Contact: Kikuya Kato

Graduate School of Biological Sciences

Nara Institute of Science and Technology

8916-5 Takayama, Ikoma, Nara 630-0101, Japan

Tel: 81-743-72-5581

Fax: 81-743-72-5589

Email: kkatob@ns.aist-nara.ac.jp,

URL: http://love2.aist-nara.ac.jp/BED/index.html.

Location/Qualifiers

1..53

/organism="Mus musculus"

/db_xref="taxon:10090"

```

/clone="BED0010931"
/clone_lib="3'-directed mouse cDNA library"
/tissue_type="brain"
/notes="Vector: pGEM-T-easy"

```

```

BASE COUNT      11 a      8 c      20 g      14 t
ORIGIN
Query Match      72.0%; Score 10.8; DB 9; Length 53;
Best Local Similarity 64.3%; Pred. No. 5.8e+04;
Matches 9; Conservative 3; Mismatches 2; Indels 0; Gaps 0;

```

```

Qy      2 GAUACGUGAAGAU 15
      |||||
Db      27 GATAGGTCAGAT 40

```

Search completed: July 6, 2003, 15:28:26
Job time : 1011.73 secs

; FILE REFERENCE: MBHB00-884-H (500/001)
; CURRENT APPLICATION NUMBER: US/09/780,929
; CURRENT FILING DATE: 2001-02-08
; PRIOR APPLICATION NUMBER: US 60/181,360
; PRIOR FILING DATE: 2000-02-08
; NUMBER OF SEQ ID NOS: 126
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 3
; LENGTH: 27
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
US-09-780-929-3

Query Match 100.0%; Score 15; DB 10; Length 27;
Best Local Similarity 100.0%; Pred. No. 90;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGAUAACGUGAAGAU 15
|||||
DB 7 AGAUAACGUGAAGAU 21

RESULT 3
US-09-780-929-126
; Sequence 126, Application US/09780929
; Patent No. US20020151693A1
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc
; APPLICANT: Breaker, Ronald
; APPLICANT: Belgelman, Leo
; TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity
; FILE REFERENCE: MBHB00-884-H (500/001)
; CURRENT APPLICATION NUMBER: US/09/780,929
; CURRENT FILING DATE: 2001-02-08
; PRIOR APPLICATION NUMBER: US 60/181,360
; PRIOR FILING DATE: 2000-02-08
; NUMBER OF SEQ ID NOS: 126
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 126
; LENGTH: 27
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
; NAME/KEY: misc_feature
; LOCATION: (1)..(6)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (21)..(27)
; OTHER INFORMATION: 2'-O-Methyl
US-09-780-929-126

Query Match 100.0%; Score 15; DB 10; Length 27;
Best Local Similarity 100.0%; Pred. No. 90;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGAUAACGUGAAGAU 15
|||||
DB 7 AGAUAACGUGAAGAU 21

RESULT 4
US-09-780-929-1
; Sequence 1, Application US/09780929
; Patent No. US20020151693A1
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc
; APPLICANT: Breaker, Ronald
; APPLICANT: Belgelman, Leo
; TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity
; FILE REFERENCE: MBHB00-884-H (500/001)

; CURRENT APPLICATION NUMBER: US/09/780,929
; CURRENT FILING DATE: 2001-02-08
; PRIOR APPLICATION NUMBER: US 60/181,360
; PRIOR FILING DATE: 2000-02-08
; NUMBER OF SEQ ID NOS: 126
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 1
; LENGTH: 28
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
; NAME/KEY: misc_feature
; LOCATION: (1)..(6)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (21)..(27)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (28)..(28)
; OTHER INFORMATION: n stands for inverted deoxybasic derivative
US-09-780-929-1

Query Match 100.0%; Score 15; DB 10; Length 28;
Best Local Similarity 100.0%; Pred. No. 91;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGAUAACGUGAAGAU 15
|||||
DB 7 AGAUAACGUGAAGAU 21

RESULT 5
US-09-780-929-2
; Sequence 2, Application US/09780929
; Patent No. US20020151693A1
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc
; APPLICANT: Breaker, Ronald
; APPLICANT: Belgelman, Leo
; TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity
; FILE REFERENCE: MBHB00-884-H (500/001)
; CURRENT APPLICATION NUMBER: US/09/780,929
; CURRENT FILING DATE: 2001-02-08
; PRIOR APPLICATION NUMBER: US 60/181,360
; PRIOR FILING DATE: 2000-02-08
; NUMBER OF SEQ ID NOS: 126
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 2
; LENGTH: 28
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
; NAME/KEY: misc_feature
; LOCATION: (1)..(5)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (21)..(27)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (28)..(28)
; OTHER INFORMATION: n stands for inverted deoxybasic derivative
US-09-780-929-2

Query Match 100.0%; Score 15; DB 10; Length 28;
Best Local Similarity 100.0%; Pred. No. 91;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGAUAACGUGAAGAU 15
|||||
DB 7 AGAUAACGUGAAGAU 21

```
RESULT 6
US-09-780-929-4
; Sequence 4, Application US/09780929
; Patent No. US20020151693A1
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc
; APPLICANT: Breaker, Ronald
; APPLICANT: Beigelman, Leo
; TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity
; FILE REFERENCE: MBH00-884-H (500/001)
; CURRENT APPLICATION NUMBER: US/09/780,929
; CURRENT FILING DATE: 2001-02-08
; PRIOR APPLICATION NUMBER: US 60/181,360
; PRIOR FILING DATE: 2000-02-08
; NUMBER OF SEQ ID NOS: 126
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 4
; LENGTH: 28
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
; NAME/KEY: misc_feature
; LOCATION: (1)..(5)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (22)..(27)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (28)..(28)
; OTHER INFORMATION: n stands for inverted deoxybasic derivative
US-09-780-929-4
Query Match 100.0%; Score 15; DB 10; Length 28;
Best Local Similarity 100.0%; Pred. No. 91;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 AGAUAACGUGAAGAU 15
Db 7 AGAUAACGUGAAGAU 21
RESULT 7
US-09-780-929-5
; Sequence 5, Application US/09780929
; Patent No. US20020151693A1
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc
; APPLICANT: Breaker, Ronald
; APPLICANT: Beigelman, Leo
; TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity
; FILE REFERENCE: MBH00-884-H (500/001)
; CURRENT APPLICATION NUMBER: US/09/780,929
; CURRENT FILING DATE: 2001-02-08
; PRIOR APPLICATION NUMBER: US 60/181,360
; PRIOR FILING DATE: 2000-02-08
; NUMBER OF SEQ ID NOS: 126
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 5
; LENGTH: 28
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
; NAME/KEY: misc_feature
; LOCATION: (1)..(5)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (23)..(27)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (28)..(28)
```

```
; OTHER INFORMATION: n stands for inverted deoxybasic derivative
US-09-780-929-5
Query Match 100.0%; Score 15; DB 10; Length 28;
Best Local Similarity 100.0%; Pred. No. 91;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 AGAUAACGUGAAGAU 15
Db 7 AGAUAACGUGAAGAU 21
RESULT 8
US-09-780-929-6
; Sequence 6, Application US/09780929
; Patent No. US20020151693A1
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc
; APPLICANT: Breaker, Ronald
; APPLICANT: Beigelman, Leo
; TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity
; FILE REFERENCE: MBH00-884-H (500/001)
; CURRENT APPLICATION NUMBER: US/09/780,929
; CURRENT FILING DATE: 2001-02-08
; PRIOR APPLICATION NUMBER: US 60/181,360
; PRIOR FILING DATE: 2000-02-08
; NUMBER OF SEQ ID NOS: 126
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 6
; LENGTH: 28
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
; NAME/KEY: misc_feature
; LOCATION: (1)..(4)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (24)..(27)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (28)..(28)
; OTHER INFORMATION: n stands for inverted deoxybasic derivative
US-09-780-929-6
Query Match 100.0%; Score 15; DB 10; Length 28;
Best Local Similarity 100.0%; Pred. No. 91;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 AGAUAACGUGAAGAU 15
Db 7 AGAUAACGUGAAGAU 21
RESULT 9
US-09-780-929-7
; Sequence 7, Application US/09780929
; Patent No. US20020151693A1
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc
; APPLICANT: Breaker, Ronald
; APPLICANT: Beigelman, Leo
; TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity
; FILE REFERENCE: MBH00-884-H (500/001)
; CURRENT APPLICATION NUMBER: US/09/780,929
; CURRENT FILING DATE: 2001-02-08
; PRIOR APPLICATION NUMBER: US 60/181,360
; PRIOR FILING DATE: 2000-02-08
; NUMBER OF SEQ ID NOS: 126
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 7
; LENGTH: 28
; TYPE: RNA
```

```
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
; NAME/KEY: misc_feature
; LOCATION: (1)..(4)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (23)..(27)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (28)..(28)
; OTHER INFORMATION: n stands for inverted deoxyabasic derivative
US-09-780-929-7

Query Match      100.0%; Score 15; DB 10; Length 28;
Best Local Similarity 100.0%; Pred. No. 91;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGAUACGUGAAGAU 15
    |||||
Db 7 AGAUACGUGAAGAU 21

RESULT 10
US-09-780-929-8
; Sequence 8, Application US/09780929
; Patent No. US20020151693A1
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc
; APPLICANT: Breaker, Ronald
; TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity
; FILE REFERENCE: MBH00-884-H (500/001)
; CURRENT APPLICATION NUMBER: US/09/780,929
; PRIOR FILING DATE: 2001-02-08
; PRIOR APPLICATION NUMBER: US 60/181,360
; NUMBER OF SEQ ID NOS: 126
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 8
; LENGTH: 28
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
; NAME/KEY: misc_feature
; LOCATION: (1)..(6)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (20)..(27)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (28)..(28)
; OTHER INFORMATION: n stands for inverted deoxyabasic derivative
US-09-780-929-8

Query Match      100.0%; Score 15; DB 10; Length 28;
Best Local Similarity 100.0%; Pred. No. 91;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGAUACGUGAAGAU 15
    |||||
Db 7 AGAUACGUGAAGAU 21

RESULT 11
US-09-780-929-9
; Sequence 9, Application US/09780929
; Patent No. US20020151693A1
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc
; APPLICANT: Breaker, Ronald
; APPLICANT: Beigelman, Leo
```

```
; TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity
; FILE REFERENCE: MBH00-884-H (500/001)
; CURRENT APPLICATION NUMBER: US/09/780,929
; PRIOR FILING DATE: 2001-02-08
; PRIOR APPLICATION NUMBER: US 60/181,360
; NUMBER OF SEQ ID NOS: 126
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 9
; LENGTH: 28
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
; NAME/KEY: misc_feature
; LOCATION: (1)..(6)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (12)..(12)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (21)..(27)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (28)..(28)
; OTHER INFORMATION: n stands for inverted deoxyabasic derivative
US-09-780-929-9

Query Match      100.0%; Score 15; DB 10; Length 28;
Best Local Similarity 100.0%; Pred. No. 91;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGAUACGUGAAGAU 15
    |||||
Db 7 AGAUACGUGAAGAU 21

RESULT 12
US-09-780-929-10
; Sequence 10, Application US/09780929
; Patent No. US20020151693A1
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc
; APPLICANT: Breaker, Ronald
; APPLICANT: Beigelman, Leo
; TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity
; FILE REFERENCE: MBH00-884-H (500/001)
; CURRENT APPLICATION NUMBER: US/09/780,929
; PRIOR FILING DATE: 2001-02-08
; PRIOR APPLICATION NUMBER: US 60/181,360
; NUMBER OF SEQ ID NOS: 126
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 10
; LENGTH: 28
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
; NAME/KEY: misc_feature
; LOCATION: (1)..(7)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (21)..(27)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (28)..(28)
; OTHER INFORMATION: n stands for inverted deoxyabasic derivative
US-09-780-929-10

Query Match      100.0%; Score 15; DB 10; Length 28;
Best Local Similarity 100.0%; Pred. No. 91;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

Qy 1 AGAUACGUGAAGAU 15
|||||
Db 7 AGAUACGUGAAGAU 21

RESULT 13

US-09-780-929-11
; Sequence 11, Application US/09780929
; Patent No. US20020151693A1
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc
; APPLICANT: Breaker, Ronald
; APPLICANT: Beigelman, Leo
; TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity
; FILE REFERENCE: MBH00-884-H (500/001)
; CURRENT APPLICATION NUMBER: US/09/780,929
; CURRENT FILING DATE: 2001-02-08
; PRIOR APPLICATION NUMBER: US 60/181,360
; PRIOR FILING DATE: 2000-02-08
; NUMBER OF SEQ ID NOS: 126
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 11
; LENGTH: 28
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
; NAME/KEY: misc_feature
; LOCATION: (1)..(6)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (21)..(27)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (28)..(28)
; OTHER INFORMATION: n stands for inverted deoxyabasic derivative
US-09-780-929-11

Query Match 100.0%; Score 15; DB 10; Length 28;
Best Local Similarity 100.0%; Pred. No. 91;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGAUACGUGAAGAU 15
|||||
Db 7 AGAUACGUGAAGAU 21

RESULT 14

US-09-780-929-12
; Sequence 12, Application US/09780929
; Patent No. US20020151693A1
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc
; APPLICANT: Breaker, Ronald
; APPLICANT: Beigelman, Leo
; TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity
; FILE REFERENCE: MBH00-884-H (500/001)
; CURRENT APPLICATION NUMBER: US/09/780,929
; CURRENT FILING DATE: 2001-02-08
; PRIOR APPLICATION NUMBER: US 60/181,360
; PRIOR FILING DATE: 2000-02-08
; NUMBER OF SEQ ID NOS: 126
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 12
; LENGTH: 28
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
; NAME/KEY: misc_feature
; LOCATION: (1)..(6)
; OTHER INFORMATION: 2'-O-Methyl

; NAME/KEY: misc_feature
; LOCATION: (18)..(18)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (21)..(27)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (28)..(28)
; OTHER INFORMATION: n stands for inverted deoxyabasic derivative
US-09-780-929-12

Query Match 100.0%; Score 15; DB 10; Length 28;
Best Local Similarity 100.0%; Pred. No. 91;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGAUACGUGAAGAU 15
|||||
Db 7 AGAUACGUGAAGAU 21

RESULT 15

US-09-780-929-13
; Sequence 13, Application US/09780929
; Patent No. US20020151693A1
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc
; APPLICANT: Breaker, Ronald
; APPLICANT: Beigelman, Leo
; TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity
; FILE REFERENCE: MBH00-884-H (500/001)
; CURRENT APPLICATION NUMBER: US/09/780,929
; CURRENT FILING DATE: 2001-02-08
; PRIOR APPLICATION NUMBER: US 60/181,360
; PRIOR FILING DATE: 2000-02-08
; NUMBER OF SEQ ID NOS: 126
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 13
; LENGTH: 28
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
; NAME/KEY: misc_feature
; LOCATION: (1)..(6)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (17)..(17)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (21)..(27)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (28)..(28)
; OTHER INFORMATION: n stands for inverted deoxyabasic derivative
US-09-780-929-13

Query Match 100.0%; Score 15; DB 10; Length 28;
Best Local Similarity 100.0%; Pred. No. 91;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGAUACGUGAAGAU 15
|||||
Db 7 AGAUACGUGAAGAU 21

Search completed: July 6, 2003, 16:52:32
Job time : 85 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: July 6, 2003, 14:40:47 ; Search time 1625.91 Seconds
(without alignments)
231.955 Million cell updates/sec

Title: US-09-780-929-97

Perfect score: 15

Sequence: 1 agaaacgugaagau 15

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 24791104 seqs, 12571243825 residues

Total number of hits satisfying chosen parameters: 12745074

Minimum DB seq length: 0

Maximum DB seq length: 60

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

Pending_Patents_NA_Main:*

- 1: /cgn2_6/ptodata/1/pna/PCTUS_COMB.seq.*
- 2: /cgn2_6/ptodata/1/pna/US06_COMB.seq.*
- 3: /cgn2_6/ptodata/1/pna/US07_COMB.seq.*
- 4: /cgn2_6/ptodata/1/pna/US080_COMB.seq.*
- 5: /cgn2_6/ptodata/1/pna/US081_COMB.seq.*
- 6: /cgn2_6/ptodata/1/pna/US082_COMB.seq.*
- 7: /cgn2_6/ptodata/1/pna/US083_COMB.seq.*
- 8: /cgn2_6/ptodata/1/pna/US084_COMB.seq.*
- 9: /cgn2_6/ptodata/1/pna/US085_COMB.seq.*
- 10: /cgn2_6/ptodata/1/pna/US086_COMB.seq.*
- 11: /cgn2_6/ptodata/1/pna/US087_COMB.seq.*
- 12: /cgn2_6/ptodata/1/pna/US088_COMB.seq.*
- 13: /cgn2_6/ptodata/1/pna/US089_COMB.seq.*
- 14: /cgn2_6/ptodata/1/pna/US090_COMB.seq.*
- 15: /cgn2_6/ptodata/1/pna/US091_COMB.seq.*
- 16: /cgn2_6/ptodata/1/pna/US092_COMB.seq.*
- 17: /cgn2_6/ptodata/1/pna/US093_COMB.seq.*
- 18: /cgn2_6/ptodata/1/pna/US094_COMB.seq.*
- 19: /cgn2_6/ptodata/1/pna/US095A_COMB.seq.*
- 20: /cgn2_6/ptodata/1/pna/US095B_COMB.seq.*
- 21: /cgn2_6/ptodata/1/pna/US095C_COMB.seq.*
- 22: /cgn2_6/ptodata/1/pna/US095D_COMB.seq.*
- 23: /cgn2_6/ptodata/1/pna/US096A_COMB.seq.*
- 24: /cgn2_6/ptodata/1/pna/US096B_COMB.seq.*
- 25: /cgn2_6/ptodata/1/pna/US096C_COMB.seq.*
- 26: /cgn2_6/ptodata/1/pna/US096D_COMB.seq.*
- 27: /cgn2_6/ptodata/1/pna/US096E_COMB.seq.*
- 28: /cgn2_6/ptodata/1/pna/US097A_COMB.seq.*
- 29: /cgn2_6/ptodata/1/pna/US097B_COMB.seq.*
- 30: /cgn2_6/ptodata/1/pna/US097C_COMB.seq.*
- 31: /cgn2_6/ptodata/1/pna/US098A_COMB.seq.*
- 32: /cgn2_6/ptodata/1/pna/US098B_COMB.seq.*
- 33: /cgn2_6/ptodata/1/pna/US098C_COMB.seq.*
- 34: /cgn2_6/ptodata/1/pna/US099A_COMB.seq.*
- 35: /cgn2_6/ptodata/1/pna/US099B_COMB.seq.*
- 36: /cgn2_6/ptodata/1/pna/US099C_COMB.seq.*
- 37: /cgn2_6/ptodata/1/pna/US099D_COMB.seq.*
- 38: /cgn2_6/ptodata/1/pna/US100A_COMB.seq.*
- 39: /cgn2_6/ptodata/1/pna/US100B_COMB.seq.*
- 40: /cgn2_6/ptodata/1/pna/US101A_COMB.seq.*
- 41: /cgn2_6/ptodata/1/pna/US101B_COMB.seq.*
- 42: /cgn2_6/ptodata/1/pna/US102A_COMB.seq.*
- 43: /cgn2_6/ptodata/1/pna/US102B_COMB.seq.*

- 44: /cgn2_6/ptodata/1/pna/US6000_COMB.seq.*
- 45: /cgn2_6/ptodata/1/pna/US6001_COMB.seq.*
- 46: /cgn2_6/ptodata/1/pna/US6002_COMB.seq.*
- 47: /cgn2_6/ptodata/1/pna/US6003_COMB.seq.*
- 48: /cgn2_6/ptodata/1/pna/US6004_COMB.seq.*
- 49: /cgn2_6/ptodata/1/pna/US6005_COMB.seq.*
- 50: /cgn2_6/ptodata/1/pna/US6006_COMB.seq.*
- 51: /cgn2_6/ptodata/1/pna/US6007_COMB.seq.*
- 52: /cgn2_6/ptodata/1/pna/US6008_COMB.seq.*
- 53: /cgn2_6/ptodata/1/pna/US6009_COMB.seq.*
- 54: /cgn2_6/ptodata/1/pna/US6010_COMB.seq.*
- 55: /cgn2_6/ptodata/1/pna/US6011_COMB.seq.*
- 56: /cgn2_6/ptodata/1/pna/US6012_COMB.seq.*
- 57: /cgn2_6/ptodata/1/pna/US6013_COMB.seq.*
- 58: /cgn2_6/ptodata/1/pna/US6014_COMB.seq.*
- 59: /cgn2_6/ptodata/1/pna/US6015_COMB.seq.*
- 60: /cgn2_6/ptodata/1/pna/US6016_COMB.seq.*
- 61: /cgn2_6/ptodata/1/pna/US6017_COMB.seq.*
- 62: /cgn2_6/ptodata/1/pna/US6018_COMB.seq.*
- 63: /cgn2_6/ptodata/1/pna/US6019_COMB.seq.*
- 64: /cgn2_6/ptodata/1/pna/US6020_COMB.seq.*
- 65: /cgn2_6/ptodata/1/pna/US6021_COMB.seq.*
- 66: /cgn2_6/ptodata/1/pna/US6022_COMB.seq.*
- 67: /cgn2_6/ptodata/1/pna/US6023_COMB.seq.*
- 68: /cgn2_6/ptodata/1/pna/US6024_COMB.seq.*
- 69: /cgn2_6/ptodata/1/pna/US6025_COMB.seq.*
- 70: /cgn2_6/ptodata/1/pna/US6026_COMB.seq.*
- 71: /cgn2_6/ptodata/1/pna/US6027_COMB.seq.*
- 72: /cgn2_6/ptodata/1/pna/US6028_COMB.seq.*
- 73: /cgn2_6/ptodata/1/pna/US6029_COMB.seq.*
- 74: /cgn2_6/ptodata/1/pna/US6030_COMB.seq.*
- 75: /cgn2_6/ptodata/1/pna/US6031_COMB.seq.*
- 76: /cgn2_6/ptodata/1/pna/US6032_COMB.seq.*
- 77: /cgn2_6/ptodata/1/pna/US6033_COMB.seq.*
- 78: /cgn2_6/ptodata/1/pna/US6034_COMB.seq.*
- 79: /cgn2_6/ptodata/1/pna/US6035_COMB.seq.*
- 80: /cgn2_6/ptodata/1/pna/US6036_COMB.seq.*
- 81: /cgn2_6/ptodata/1/pna/US6037_COMB.seq.*
- 82: /cgn2_6/ptodata/1/pna/US6038_COMB.seq.*
- 83: /cgn2_6/ptodata/1/pna/US6039_COMB.seq.*
- 84: /cgn2_6/ptodata/1/pna/US6040_COMB.seq.*
- 85: /cgn2_6/ptodata/1/pna/US6041_COMB.seq.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	15	100.0	15	30	US-09-780-929-97
2	15	100.0	27	30	US-09-780-929-3
3	15	100.0	27	30	US-09-780-929-126
4	15	100.0	28	30	US-09-780-929-1
5	15	100.0	28	30	US-09-780-929-2
6	15	100.0	28	30	US-09-780-929-4
7	15	100.0	28	30	US-09-780-929-5
8	15	100.0	28	30	US-09-780-929-6
9	15	100.0	28	30	US-09-780-929-7
10	15	100.0	28	30	US-09-780-929-8
11	15	100.0	28	30	US-09-780-929-9
12	15	100.0	28	30	US-09-780-929-10
13	15	100.0	28	30	US-09-780-929-11
14	15	100.0	28	30	US-09-780-929-12
15	15	100.0	28	30	US-09-780-929-13
16	15	100.0	28	30	US-09-780-929-14
17	15	100.0	28	30	US-09-780-929-15
18	15	100.0	28	30	US-09-780-929-16
19	15	100.0	28	30	US-09-780-929-17
20	15	100.0	28	30	US-09-780-929-18
21	15	100.0	28	30	US-09-780-929-19

```
22 15 100.0 28 30 US-09-780-929-20 Sequence 20, Appl
23 15 100.0 28 30 US-09-780-929-21 Sequence 21, Appl
24 15 100.0 28 30 US-09-780-929-22 Sequence 22, Appl
25 15 100.0 28 30 US-09-780-929-23 Sequence 23, Appl
26 15 100.0 28 30 US-09-780-929-24 Sequence 24, Appl
27 15 100.0 28 30 US-09-780-929-25 Sequence 25, Appl
28 15 100.0 28 30 US-09-780-929-26 Sequence 26, Appl
29 15 100.0 28 30 US-09-780-929-27 Sequence 27, Appl
30 15 100.0 28 30 US-09-780-929-28 Sequence 28, Appl
31 15 100.0 28 30 US-09-780-929-29 Sequence 29, Appl
32 15 100.0 28 30 US-09-780-929-30 Sequence 30, Appl
33 15 100.0 28 30 US-09-780-929-31 Sequence 31, Appl
34 15 100.0 28 30 US-09-780-929-32 Sequence 32, Appl
35 15 100.0 28 30 US-09-780-929-33 Sequence 33, Appl
36 15 100.0 28 30 US-09-780-929-34 Sequence 34, Appl
37 15 100.0 28 30 US-09-780-929-35 Sequence 35, Appl
38 15 100.0 28 30 US-09-780-929-36 Sequence 36, Appl
39 15 100.0 28 30 US-09-780-929-37 Sequence 37, Appl
40 15 100.0 28 30 US-09-780-929-38 Sequence 38, Appl
41 15 100.0 28 30 US-09-780-929-39 Sequence 39, Appl
42 15 100.0 28 30 US-09-780-929-40 Sequence 40, Appl
43 15 100.0 28 30 US-09-780-929-41 Sequence 41, Appl
44 15 100.0 28 30 US-09-780-929-42 Sequence 42, Appl
45 15 100.0 28 30 US-09-780-929-43 Sequence 43, Appl
```

ALIGNMENTS

```
RESULT 1
US-09-780-929-97
; Sequence 97, Application US/09780929
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc
; APPLICANT: Breaker, Ronald
; APPLICANT: Beigelman, Leo
; TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity
; FILE REFERENCE: MBH00-884-H (500/001)
; CURRENT APPLICATION NUMBER: US/09/780,929
; CURRENT FILING DATE: 2001-02-08
; PRIOR APPLICATION NUMBER: US 60/181,360
; PRIOR FILING DATE: 2000-02-08
; NUMBER OF SEQ ID NOS: 126
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 97
; LENGTH: 15
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
US-09-780-929-97
```

```
Query Match 100.0%; Score 15; DB 30; Length 15;
Best Local Similarity 100.0%; Pred. No. 3.8e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

```
Oy 1 AGAUACGUGAAGAU 15
Db 1 AGAUACGUGAAGAU 15
|||||
```

```
RESULT 2
US-09-780-929-3
; Sequence 3, Application US/09780929
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc
; APPLICANT: Breaker, Ronald
; APPLICANT: Beigelman, Leo
; TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity
; FILE REFERENCE: MBH00-884-H (500/001)
; CURRENT APPLICATION NUMBER: US/09/780,929
; CURRENT FILING DATE: 2001-02-08
; PRIOR APPLICATION NUMBER: US 60/181,360
```

```
; PRIOR FILING DATE: 2000-02-08
; NUMBER OF SEQ ID NOS: 126
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 3
; LENGTH: 27
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
US-09-780-929-3
```

```
Query Match 100.0%; Score 15; DB 30; Length 27;
Best Local Similarity 100.0%; Pred. No. 4.2e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

```
Oy 1 AGAUACGUGAAGAU 15
Db 7 AGAUACGUGAAGAU 21
|||||
```

```
RESULT 3
US-09-780-929-126
; Sequence 126, Application US/09780929
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc
; APPLICANT: Breaker, Ronald
; APPLICANT: Beigelman, Leo
; TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity
; FILE REFERENCE: MBH00-884-H (500/001)
; CURRENT APPLICATION NUMBER: US/09/780,929
; CURRENT FILING DATE: 2001-02-08
; PRIOR APPLICATION NUMBER: US 60/181,360
; PRIOR FILING DATE: 2000-02-08
; NUMBER OF SEQ ID NOS: 126
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 126
; LENGTH: 27
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
; NAME/KEY: misc.feature
; LOCATION: (1)..(6)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc.feature
; LOCATION: (21)..(27)
; OTHER INFORMATION: 2'-O-Methyl
US-09-780-929-126
```

```
Query Match 100.0%; Score 15; DB 30; Length 27;
Best Local Similarity 100.0%; Pred. No. 4.2e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

```
Oy 1 AGAUACGUGAAGAU 15
Db 7 AGAUACGUGAAGAU 21
|||||
```

```
RESULT 4
US-09-780-929-1
; Sequence 1, Application US/09780929
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc
; APPLICANT: Breaker, Ronald
; APPLICANT: Beigelman, Leo
; TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity
; FILE REFERENCE: MBH00-884-H (500/001)
; CURRENT APPLICATION NUMBER: US/09/780,929
; CURRENT FILING DATE: 2001-02-08
; PRIOR APPLICATION NUMBER: US 60/181,360
; PRIOR FILING DATE: 2000-02-08
; NUMBER OF SEQ ID NOS: 126
; SOFTWARE: PatentIn version 3.0
```

```
; SEQ ID NO 1
; LENGTH: 28
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
; NAME/KEY: misc_feature
; LOCATION: (1)..(6)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (21)..(27)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (28)..(28)
; OTHER INFORMATION: n stands for inverted deoxyabasic derivative
US-09-780-929-1

Query Match          100.0%; Score 15; DB 30; Length 28;
Best Local Similarity 100.0%; Pred. No. 4.2e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGAUAAACGUGAAGAU 15
    |||||
Db 7 AGAUAAACGUGAAGAU 21

RESULT 5
US-09-780-929-2
; Sequence 2, Application US/09780929
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc
; APPLICANT: Breaker, Ronald
; APPLICANT: Beigelman, Leo
; TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity
; FILE REFERENCE: MBH00-884-H (500/001)
; CURRENT APPLICATION NUMBER: US/09/780,929
; CURRENT FILING DATE: 2001-02-08
; PRIOR APPLICATION NUMBER: US 60/181,360
; PRIOR FILING DATE: 2000-02-08
; NUMBER OF SEQ ID NOS: 126
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 2
; LENGTH: 28
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
; NAME/KEY: misc_feature
; LOCATION: (1)..(5)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (21)..(27)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (28)..(28)
; OTHER INFORMATION: n stands for inverted deoxyabasic derivative
US-09-780-929-2

Query Match          100.0%; Score 15; DB 30; Length 28;
Best Local Similarity 100.0%; Pred. No. 4.2e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGAUAAACGUGAAGAU 15
    |||||
Db 7 AGAUAAACGUGAAGAU 21

RESULT 6
US-09-780-929-4
; Sequence 4, Application US/09780929
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc
; APPLICANT: Breaker, Ronald
```

```
; APPLICANT: Beigelman, Leo
; TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity
; FILE REFERENCE: MBH00-884-H (500/001)
; CURRENT APPLICATION NUMBER: US/09/780,929
; CURRENT FILING DATE: 2001-02-08
; PRIOR APPLICATION NUMBER: US 60/181,360
; PRIOR FILING DATE: 2000-02-08
; NUMBER OF SEQ ID NOS: 126
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 4
; LENGTH: 28
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
; NAME/KEY: misc_feature
; LOCATION: (1)..(5)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (22)..(27)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (28)..(28)
; OTHER INFORMATION: n stands for inverted deoxyabasic derivative
US-09-780-929-4

Query Match          100.0%; Score 15; DB 30; Length 28;
Best Local Similarity 100.0%; Pred. No. 4.2e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGAUAAACGUGAAGAU 15
    |||||
Db 7 AGAUAAACGUGAAGAU 21

RESULT 7
US-09-780-929-5
; Sequence 5, Application US/09780929
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc
; APPLICANT: Breaker, Ronald
; APPLICANT: Beigelman, Leo
; TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity
; FILE REFERENCE: MBH00-884-H (500/001)
; CURRENT APPLICATION NUMBER: US/09/780,929
; CURRENT FILING DATE: 2001-02-08
; PRIOR APPLICATION NUMBER: US 60/181,360
; PRIOR FILING DATE: 2000-02-08
; NUMBER OF SEQ ID NOS: 126
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 5
; LENGTH: 28
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
; NAME/KEY: misc_feature
; LOCATION: (1)..(5)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (23)..(27)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (28)..(28)
; OTHER INFORMATION: n stands for inverted deoxyabasic derivative
US-09-780-929-5

Query Match          100.0%; Score 15; DB 30; Length 28;
Best Local Similarity 100.0%; Pred. No. 4.2e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGAUAAACGUGAAGAU 15
    |||||
```

```
Db      7 AGAUAACGUGAAGAU 21

RESULT 8
US-09-780-929-6
; Sequence 6, Application US/09780929
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc
; APPLICANT: Breaker, Ronald
; APPLICANT: Beigelman, Leo
; TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity
; FILE REFERENCE: MBH00-884-H (500/001)
; CURRENT APPLICATION NUMBER: US/09/780,929
; CURRENT FILING DATE: 2001-02-08
; PRIOR APPLICATION NUMBER: US 60/181,360
; PRIOR FILING DATE: 2000-02-08
; NUMBER OF SEQ ID NOS: 126
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 6
; LENGTH: 28
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
; NAME/KEY: misc_feature
; LOCATION: (1)..(4)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (24)..(27)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (28)..(28)
; OTHER INFORMATION: n stands for inverted deoxybasic derivative
US-09-780-929-6

Query Match      100.0%; Score 15; DB 30; Length 28;
Best Local Similarity 100.0%; Pred. No. 4.2e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 AGAUAACGUGAAGAU 15
        |||||
Db      7 AGAUAACGUGAAGAU 21

RESULT 10
US-09-780-929-8
; Sequence 8, Application US/09780929
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc
; APPLICANT: Breaker, Ronald
; APPLICANT: Beigelman, Leo
; TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity
; FILE REFERENCE: MBH00-884-H (500/001)
; CURRENT APPLICATION NUMBER: US/09/780,929
; CURRENT FILING DATE: 2001-02-08
; PRIOR APPLICATION NUMBER: US 60/181,360
; PRIOR FILING DATE: 2000-02-08
; NUMBER OF SEQ ID NOS: 126
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 8
; LENGTH: 28
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
; NAME/KEY: misc_feature
; LOCATION: (1)..(6)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (20)..(27)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (28)..(28)
; OTHER INFORMATION: n stands for inverted deoxybasic derivative
US-09-780-929-8

Query Match      100.0%; Score 15; DB 30; Length 28;
Best Local Similarity 100.0%; Pred. No. 4.2e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 AGAUAACGUGAAGAU 15
        |||||
Db      7 AGAUAACGUGAAGAU 21

RESULT 11
US-09-780-929-9
; Sequence 9, Application US/09780929
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc
; APPLICANT: Breaker, Ronald
; APPLICANT: Beigelman, Leo
; TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity
; FILE REFERENCE: MBH00-884-H (500/001)
; CURRENT APPLICATION NUMBER: US/09/780,929
; CURRENT FILING DATE: 2001-02-08
; PRIOR APPLICATION NUMBER: US 60/181,360
; PRIOR FILING DATE: 2000-02-08
; NUMBER OF SEQ ID NOS: 126
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 9
; LENGTH: 28
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
; NAME/KEY: misc_feature
; LOCATION: (1)..(4)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (23)..(27)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (28)..(28)
```



```
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
; NAME/KEY: misc_feature
; LOCATION: (1)..(6)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (12)..(12)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (21)..(27)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (28)..(28)
; OTHER INFORMATION: n stands for inverted deoxybasic derivative
US-09-780-929-9
Query Match          100.0%; Score 15; DB 30; Length 28;
Best Local Similarity 100.0%; Pred. No. 4.2e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGAUACGUGAAGAU 15
   |||||
Db 7 AGAUACGUGAAGAU 21

RESULT 12
US-09-780-929-10
; Sequence 10, Application US/09780929
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc
; APPLICANT: Breaker, Ronald
; TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity
; FILE REFERENCE: MBH00-884-H (500/001)
; CURRENT APPLICATION NUMBER: US/09/780,929
; CURRENT FILING DATE: 2001-02-08
; PRIOR APPLICATION NUMBER: US 60/181,360
; PRIOR FILING DATE: 2000-02-08
; NUMBER OF SEQ ID NOS: 126
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 10
; LENGTH: 28
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
; NAME/KEY: misc_feature
; LOCATION: (1)..(7)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (21)..(27)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (28)..(28)
; OTHER INFORMATION: n stands for inverted deoxybasic derivative
US-09-780-929-10
Query Match          100.0%; Score 15; DB 30; Length 28;
Best Local Similarity 100.0%; Pred. No. 4.2e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGAUACGUGAAGAU 15
   |||||
Db 7 AGAUACGUGAAGAU 21

RESULT 13
US-09-780-929-11
; Sequence 11, Application US/09780929
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc
; APPLICANT: Breaker, Ronald
; TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity
```

```
; FILE REFERENCE: MBH00-884-H (500/001)
; CURRENT APPLICATION NUMBER: US/09/780,929
; CURRENT FILING DATE: 2001-02-08
; PRIOR APPLICATION NUMBER: US 60/181,360
; PRIOR FILING DATE: 2000-02-08
; NUMBER OF SEQ ID NOS: 126
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 11
; LENGTH: 28
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
; NAME/KEY: misc_feature
; LOCATION: (1)..(6)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (21)..(27)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (28)..(28)
; OTHER INFORMATION: n stands for inverted deoxybasic derivative
US-09-780-929-11
Query Match          100.0%; Score 15; DB 30; Length 28;
Best Local Similarity 100.0%; Pred. No. 4.2e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGAUACGUGAAGAU 15
   |||||
Db 7 AGAUACGUGAAGAU 21

RESULT 14
US-09-780-929-12
; Sequence 12, Application US/09780929
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc
; APPLICANT: Breaker, Ronald
; TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity
; FILE REFERENCE: MBH00-884-H (500/001)
; CURRENT APPLICATION NUMBER: US/09/780,929
; CURRENT FILING DATE: 2001-02-08
; PRIOR APPLICATION NUMBER: US 60/181,360
; PRIOR FILING DATE: 2000-02-08
; NUMBER OF SEQ ID NOS: 126
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 12
; LENGTH: 28
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
; NAME/KEY: misc_feature
; LOCATION: (1)..(6)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (18)..(18)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (21)..(27)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (28)..(28)
; OTHER INFORMATION: n stands for inverted deoxybasic derivative
US-09-780-929-12
Query Match          100.0%; Score 15; DB 30; Length 28;
Best Local Similarity 100.0%; Pred. No. 4.2e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGAUACGUGAAGAU 15
   |||||
Db 7 AGAUACGUGAAGAU 21
```

```

Db      7 AGAUACGUGAAGAU 21
|||||
RESULT 15
US-09-780-929-13
; Sequence 13, Application US/09780929
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc
; APPLICANT: Breaker, Ronald
; APPLICANT: Beigelman, Leo
; TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity
; FILE REFERENCE: MBH00-884-H (500/001)
; CURRENT APPLICATION NUMBER: US/09/780,929
; CURRENT FILING DATE: 2001-02-08
; PRIOR APPLICATION NUMBER: US 60/181,360
; PRIOR FILING DATE: 2000-02-08
; NUMBER OF SEQ ID NOS: 126
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 13
; LENGTH: 28
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)..(6)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (17)..(17)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (21)..(27)
; OTHER INFORMATION: 2'-O-Methyl
; NAME/KEY: misc_feature
; LOCATION: (28)..(28)
; OTHER INFORMATION: n stands for inverted deoxyabasic derivative
US-09-780-929-13
Query Match      100.0%; Score 15; DB 30; Length 28;
Best Local Similarity 100.0%; Pred. No. 4.2e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy      1 AGAUACGUGAAGAU 15
|||||
Db      7 AGAUACGUGAAGAU 21

```

Search completed: July 6, 2003, 16:29:53
Job time : 1625.91 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2003 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: July 6, 2003, 14:36:16 ; Search time 591.818 Seconds
(without alignments)
885.154 Million cell updates/sec

Title: US-09-780-929-98

Perfect score: 18

Sequence: 1 aaugccuauccggugcga 18

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 2054640 seqs, 14551402878 residues

Total number of hits satisfying chosen parameters: 897812

Minimum DB seq length: 0

Maximum DB seq length: 60

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :

GenEmbl.*

1: gb.ba.*

2: gb.htg.*

3: gb.in.*

4: gb.om.*

5: gb.ov.*

6: gb.pat.*

7: gb.ph.*

8: gb.pl.*

9: gb.pr.*

10: gb.ro.*

11: gb.sts.*

12: gb.sy.*

13: gb.un.*

14: gb.vi.*

15: em.ba.*

16: em.fun.*

17: em.hum.*

18: em.in.*

19: em.mu.*

20: em.om.*

21: em.or.*

22: em.ov.*

23: em.pat.*

24: em.ph.*

25: em.pl.*

26: em.ro.*

27: em.sts.*

28: em.un.*

29: em.vi.*

30: em.htg.hum.*

31: em.htg.inv.*

32: em.htg.other.*

33: em.htg.mus.*

34: em.htg.pln.*

35: em.htg.rod.*

36: em.htg.mam.*

37: em.htg.vrt.*

38: em.sy.*

39: em.htgo.hum.*

40: em.htgo.mus.*

41: em.htgo.other.*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Match	Length	DB	ID	Description
1	18	100.0	18	6	AX214296	Sequence
2	18	100.0	29	6	AX214316	Sequence
3	12.8	71.1	20	6	AX295903	Sequence
4	12.8	71.1	24	6	AX291270	Sequence
5	12.2	67.8	20	6	AX293899	Sequence
6	12.2	67.8	24	6	AX289266	Sequence
7	12.2	67.8	29	6	AR198859	Sequence
8	12.2	66.7	22	6	AX297915	Sequence
9	11.8	65.6	20	6	AX293092	Sequence
10	11.8	65.6	20	6	E30812	Novel prote
11	11.8	65.6	22	6	AX019596	Sequence
12	11.8	65.6	24	6	AX288459	Sequence
13	11.8	65.6	24	6	AX447444	Sequence
14	11.8	65.6	30	6	A29209	DNA probe f
15	11.8	65.6	30	6	A29212	Oligonucleo
16	11.8	65.6	40	6	AR178716	Sequence
17	11.8	65.6	40	6	AR205421	Sequence
18	11.6	64.4	22	6	AX166857	Sequence
19	11.6	64.4	24	6	AX166856	Sequence
20	11.6	64.4	25	6	AX166855	Sequence
21	11.6	64.4	26	6	I22031	Sequence 5
22	11.6	64.4	29	6	AR023454	Sequence
23	11.6	64.4	29	6	E46848	Derivative
24	11.6	64.4	30	6	AR023453	Sequence
25	11.6	64.4	30	6	AR092568	Sequence
26	11.6	64.4	30	6	AR139964	Sequence
27	11.6	64.4	30	6	AR140283	Sequence
28	11.6	64.4	30	6	AR140561	Sequence
29	11.6	64.4	30	6	E46847	Derivative
30	11.6	64.4	41	6	A63966	Sequence 10
31	11.6	64.4	41	6	AR091476	Sequence
32	11.6	64.4	46	6	AR170886	Sequence
33	11.6	64.4	47	6	AX378262	Sequence
34	11.4	63.3	17	6	AR057682	Sequence
35	11.4	63.3	17	6	AR057773	Sequence
36	11.4	63.3	17	6	AR115440	Sequence
37	11.4	63.3	17	6	AR115531	Sequence
38	11.4	63.3	21	6	AR167007	Sequence
39	11.4	63.3	21	6	AR210662	Sequence
40	11.4	63.3	24	6	AX447141	Sequence
41	11.4	63.3	25	6	AR097506	Sequence
42	11.4	63.3	25	6	AR139820	Sequence
43	11.4	63.3	25	6	AR140127	Sequence
44	11.4	63.3	25	6	AR142844	Sequence
45	11.4	63.3	31	6	AX425973	Sequence

ALIGNMENTS

RESULT 1	AX214296	AX214296	18 bp	mRNA	linear	PAT 06-SEP-2001
LOCUS	Sequence 109 from Patent WO0159102.					
DEFINITION	AX214296					
ACCESSION	AX214296					
VERSION	AX214296.1	GI:15524373				
KEYWORDS	Synthetic construct.					
SOURCE	Synthetic construct.					
ORGANISM	artificial sequences.					
REFERENCE	1 (bases 1 to 18)					
AUTHORS	Breaker R. and Emilsson G.					
TITLE	Nucleozymes with endonuclease activity					
JOURNAL	Patent: WO 0159102-A 109 16-AUG-2001;					
	RIBOZYME PHARMACEUTICALS, INC. (US) ; Yale University (US)					

```
FEATURES
  source
    Location/Qualifiers
      1..18
        /organism="synthetic construct"
        /db_xref="taxon:32630"
        /note="Nucleic Acid"
      4 a 4 c 6 g 4 t
BASE COUNT
  4 a 4 c 6 g 4 t
ORIGIN
  Query Match
    Best Local Similarity 100.0%; Score 18; DB 6; Length 18;
    Matches 14; Conservative 4; Mismatches 0; Indels 0; Gaps 0;
  Qy
    1 AAUGGCCUAUCGGUGCGA 18
      ||:||||:|:|||||
  Db
    1 AATGGCCTATCGGTGCGA 18
      ||:||||:|:|||||
RESULT 2
  LOCUS
    AX214316 29 bp mRNA linear PAT 06-SEP-2001
  DEFINITION
    Sequence 129 from Patent WO0159102.
  ACCESSION
    AX214316
  VERSION
    AX214316.1 GI:15524393
  KEYWORDS
    synthetic construct.
  SOURCE
    synthetic construct.
    artificial sequences.
  ORGANISM
    Barany,F., Zirvi,M., Gerry,N.P., Favis,R. and Kliman,R.
  REFERENCE
    1 (bases 1 to 29)
  AUTHORS
    Breaker,R. and Emilsson,G.
  TITLE
    Nucleozymes with endonuclease activity
  JOURNAL
    Patent: WO 0159102-A 129 16-AUG-2001.
  JOURNAL
    RIBOZYME PHARMACEUTICALS, INC. (US) ; Yale University (US)
FEATURES
  source
    Location/Qualifiers
      1..29
        /organism="synthetic construct"
        /db_xref="taxon:32630"
        /note="Nucleic Acid"
      6 a 7 c 11 g 5 t
BASE COUNT
  6 a 7 c 11 g 5 t
ORIGIN
  Query Match
    Best Local Similarity 100.0%; Score 18; DB 6; Length 29;
    Matches 14; Conservative 4; Mismatches 0; Indels 0; Gaps 0;
  Qy
    1 AAUGGCCUAUCGGUGCGA 18
      ||:||||:|:|||||
  Db
    8 AATGGCCTATCGGTGCGA 25
      ||:||||:|:|||||
RESULT 3
  LOCUS
    AX295903 20 bp DNA linear PAT 21-NOV-2001
  DEFINITION
    Sequence 7665 from Patent WO0179548.
  ACCESSION
    AX295903
  VERSION
    AX295903.1 GI:17057592
  KEYWORDS
    synthetic construct.
  SOURCE
    synthetic construct.
    artificial sequences.
  ORGANISM
    Barany,F., Zirvi,M., Gerry,N.P., Favis,R. and Kliman,R.
  REFERENCE
    1
  AUTHORS
    Method of designing addressable array for detection of nucleic acid
  TITLE
    sequence differences using ligase detection reaction
  JOURNAL
    Patent: WO 0179548-A 7665 25-OCT-2001;
  JOURNAL
    CORNELL RESEARCH FOUNDATION, INC. (US)
FEATURES
  source
    Location/Qualifiers
      1..20
        /organism="synthetic construct"
        /db_xref="taxon:32630"
        /note="Hypothetical Probe Sequence"
      2 a 6 c 6 g 6 t
BASE COUNT
  2 a 6 c 6 g 6 t
ORIGIN
  Query Match
    Best Local Similarity 77.8%; Score 18; DB 6; Length 29;
    Matches 14; Conservative 4; Mismatches 0; Indels 0; Gaps 0;
  Qy
    1 AAUGGCCUAUCGGUGCGA 18
      ||:||||:|:|||||
  Db
    8 AATGGCCTATCGGTGCGA 25
      ||:||||:|:|||||
RESULT 4
  LOCUS
    AX291270 24 bp DNA linear PAT 21-NOV-2001
  DEFINITION
    Sequence 3032 from Patent WO0179548.
  ACCESSION
    AX291270
  VERSION
    AX291270.1 GI:17052953
  KEYWORDS
    synthetic construct.
  SOURCE
    synthetic construct.
    artificial sequences.
  ORGANISM
    Barany,F., Zirvi,M., Gerry,N.P., Favis,R. and Kliman,R.
  REFERENCE
    1
  AUTHORS
    Method of designing addressable array for detection of nucleic acid
  TITLE
    sequence differences using ligase detection reaction
  JOURNAL
    Patent: WO 0179548-A 3032 25-OCT-2001;
  JOURNAL
    CORNELL RESEARCH FOUNDATION, INC. (US)
FEATURES
  source
    Location/Qualifiers
      1..24
        /organism="synthetic construct"
        /db_xref="taxon:32630"
        /note="Hypothetical Probe Sequence"
      2 a 7 c 7 g 8 t
BASE COUNT
  2 a 7 c 7 g 8 t
ORIGIN
  Query Match
    Best Local Similarity 71.1%; Score 12.8; DB 6; Length 24;
    Matches 11; Conservative 3; Mismatches 2; Indels 0; Gaps 0;
  Qy
    2 AUGGCCUAUCGGUGCG 17
      | ||| :|:|:|
  Db
    2 ACGGCTTATCGGTGCG 17
      | ||| :|:|:|
RESULT 5
  LOCUS
    AX293899 20 bp DNA linear PAT 21-NOV-2001
  DEFINITION
    Sequence 5661 from Patent WO0179548.
  ACCESSION
    AX293899
  VERSION
    AX293899.1 GI:17055582
  KEYWORDS
    synthetic construct.
  SOURCE
    synthetic construct.
    artificial sequences.
  ORGANISM
    Barany,F., Zirvi,M., Gerry,N.P., Favis,R. and Kliman,R.
  REFERENCE
    1
  AUTHORS
    Method of designing addressable array for detection of nucleic acid
  TITLE
    sequence differences using ligase detection reaction
  JOURNAL
    Patent: WO 0179548-A 5661 25-OCT-2001;
  JOURNAL
    CORNELL RESEARCH FOUNDATION, INC. (US)
FEATURES
  source
    Location/Qualifiers
      1..20
        /organism="synthetic construct"
        /db_xref="taxon:32630"
        /note="Hypothetical Probe Sequence"
      7 a 6 c 4 g 3 t
BASE COUNT
  7 a 6 c 4 g 3 t
ORIGIN
  Query Match
    Best Local Similarity 67.8%; Score 12.2; DB 6; Length 20;
    Matches 11; Conservative 3; Mismatches 3; Indels 0; Gaps 0;
  Qy
    2 AUGGCCUAUCGGUGCGA 18
      |:| || |:| | :|
  Db
    3 ATGACCAATCGATGCGA 19
      |:| || |:| | :|
```

RESULT 6
AX289266
LOCUS AX289266 24 bp DNA linear PAT 21-NOV-2001
DEFINITION Sequence 1028 from Patent WO0179548.
ACCESSION AX289266
VERSION AX289266.1 GI:17050949
KEYWORDS synthetic construct.
SOURCE synthetic construct.
ORGANISM artificial sequences.
REFERENCE 1
Barany, F., Zirvi, M., Gerry, N.P., Favis, R. and Kliman, R.
AUTHORS Method of designing addressable array for detection of nucleic acid
TITLE sequence differences using ligase detection reaction
JOURNAL Patent: WO 0179548-A 1028 25-OCT-2001;
CORNELL RESEARCH FOUNDATION, INC. (US)
FEATURES Location/Qualifiers
source 1..24
/organism="synthetic construct"
/db_xref="taxon:32630"
/note="Hypothetical Probe Sequence"
BASE COUNT 7 a 7 c 5 g 5 t
ORIGIN

Query Match 67.8%; Score 12.2; DB 6; Length 24;
Best Local Similarity 64.7%; Pred. No. 5.5e+04;
Matches 11; Conservative 3; Mismatches 3; Indels 0; Gaps 0;

QY 2 AUGGCCUAUCGGUGCGA 18
I:|||||:|||||
Db 7 ATGACCAATCGATGCGA 23

RESULT 7
AR198859/c
LOCUS AR198859 29 bp DNA linear PAT 20-APR-2002
DEFINITION Sequence 144 from patent US 6355411.
ACCESSION AR198859
VERSION AR198859.1 GI:20248933
KEYWORDS unknown.
SOURCE unknown.
ORGANISM unclassified.
REFERENCE 1 (bases 1 to 29)
AUTHORS Ausubel, F., Goodman, H.M., Rahme, L.G., Mahajan-Miklos, S., Tan, M.-W.,
Cao, H., Drenth, E. and Tsongalis, J.
TITLE Virulence-associated nucleic acid sequences and uses thereof
JOURNAL Patent: US 6355411-A 144 12-MAR-2002;
FEATURES Location/Qualifiers
source 1..29
/organism="unknown"
BASE COUNT 6 a 10 c 8 g 5 t
ORIGIN

Query Match 67.8%; Score 12.2; DB 6; Length 29;
Best Local Similarity 64.7%; Pred. No. 5.5e+04;
Matches 11; Conservative 3; Mismatches 3; Indels 0; Gaps 0;

QY 1 AAUGGCCUAUCGGUGCG 17
I:|||||:|||||
Db 19 AACGGCGTATCGTTCG 3

RESULT 8
AX297915
LOCUS AX297915 22 bp DNA linear PAT 26-NOV-2001
DEFINITION Sequence 4 from Patent WO0183757.
ACCESSION AX297915
VERSION AX297915.1 GI:17128036
KEYWORDS TT virus.
SOURCE

ORGANISM TT virus
Viruses; ssDNA viruses; unclassified ssDNA viruses.
REFERENCE 1
AUTHORS Ott, C. and Komurian-Pradel, F.
TITLE TT virus polypeptide, nucleic acid coding for said polypeptide and
uses
JOURNAL Patent: WO 0183757-A 4 08-NOV-2001;
BIO.MERIEUX (FR)
FEATURES Location/Qualifiers
source 1..22
/organism="TT virus"
/db_xref="taxon:68887"
BASE COUNT 4 a 5 c 8 g 4 t 1 others
ORIGIN

Query Match 66.7%; Score 12; DB 6; Length 22;
Best Local Similarity 57.1%; Pred. No. 7.1e+04;
Matches 8; Conservative 5; Mismatches 1; Indels 0; Gaps 0;

QY 2 AUGGCCUAUCGGUG 15
I:|||||:|:
Db 8 ATGGCCTATGGSTG 21

RESULT 9
AX293092
LOCUS AX293092 20 bp DNA linear PAT 21-NOV-2001
DEFINITION Sequence 4854 from Patent WO0179548.
ACCESSION AX293092
VERSION AX293092.1 GI:17054775
KEYWORDS synthetic construct.
SOURCE synthetic construct.
ORGANISM artificial sequences.
REFERENCE 1
AUTHORS Barany, F., Zirvi, M., Gerry, N.P., Favis, R. and Kliman, R.
TITLE Method of designing addressable array for detection of nucleic acid
sequence differences using ligase detection reaction
JOURNAL Patent: WO 0179548-A 4854 25-OCT-2001;
CORNELL RESEARCH FOUNDATION, INC. (US)
FEATURES Location/Qualifiers
source 1..20
/organism="synthetic construct"
/db_xref="taxon:32630"
/note="Hypothetical Probe Sequence"
BASE COUNT 1 a 6 c 6 g 7 t
ORIGIN

Query Match 65.6%; Score 11.8; DB 6; Length 20;
Best Local Similarity 60.0%; Pred. No. 9.2e+04;
Matches 9; Conservative 4; Mismatches 2; Indels 0; Gaps 0;

QY 3 UGGCCUAUCGGUGCG 17
I:|||||:|:
Db 6 TGCCCTATCTGTGCG 20

RESULT 10
E30812/c
LOCUS E30812 20 bp DNA linear PAT 18-JUN-2001
DEFINITION Novel protein participating in differentiation of cranial nerve
tissue cell.
ACCESSION E30812
VERSION E30812.1 GI:13017242
KEYWORDS JP 1999318468-A/6.
SOURCE unidentified.
ORGANISM unclassified.
REFERENCE 1 (bases 1 to 20)
AUTHORS Shigeru, N. and Hirofumi, S.
TITLE Novel protein participating in differentiation of cranial nerve
tissue cell
JOURNAL Patent: JP 1999318468-A 6 24-NOV-1999;

THE KANAGAWA ACADEMY OF SCIENCE, MEIJI MILK PROD CO LTD
OS Unidentified
PN JP 199318468-A/6
PD 24-NOV-1999
PF 15-MAY-1998 JP 1998152027
PR SHIGERU NOGUCHI, HIROFUMI SUEMORI
PI C12N15/09, A01K67/027, A61K38/00, A61K48/00, C07K14/47, C07K16/18,
PC C12N1/19,
PC C12N1/21, C12N5/10, C12P21/02, C12Q1/68, (C12N5/10, C12R1/91), PC
C12N15/00,
PC A61K37/02, C12N5/00, (C12N5/00, C12R1/91)
CC Strandedness: Single;
FH Key Location/Qualifiers
FT source 1..20 /organism="Unidentified".
FEATURES source Location/Qualifiers
1..20 /organism="unidentified"
/db_xref="taxon:32644"
BASE COUNT 7 a 7 c 4 g 2 t
ORIGIN
Query Match 65.6%; Score 11.8; DB 6; Length 20;
Best Local Similarity 60.0%; Pred. No. 9.2e+04;
Matches 9; Conservative 4; Mismatches 2; Indels 0; Gaps 0;
Qy 3 UGGCCUAUCGGUGCG 17
Db 20 TGGTCTCTCGTGGC 6
RESULT 11
AX019596
LOCUS
DEFINITION Sequence 50 from Patent WO938964.
AX019596
ACCESSION AX019596
VERSION AX019596.1 GI:10043510
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct
artificial sequences.
REFERENCE 1
AUTHORS Keith, W.N.
TITLE Promoter regions of the mouse and human telomerase rna component
genes
JOURNAL Patent: WO 938964-A 50 05-AUG-1999;
KEITH WILLIAM NICOL (GB); CANCER RES CAMPAIGN TECH (GB)
FEATURES source Location/Qualifiers
1..22 /organism="synthetic construct"
/db_xref="taxon:32630"
/note="Oligonucleotide"
BASE COUNT 2 a 7 c 8 g 5 t
ORIGIN
Query Match 65.6%; Score 11.8; DB 6; Length 22;
Best Local Similarity 66.7%; Pred. No. 9.2e+04;
Matches 10; Conservative 3; Mismatches 2; Indels 0; Gaps 0;
Qy 3 UGGCCUAUCGGUGCG 17
Db 3 TGGCCAAATCCGTGGC 17
RESULT 12
AX288459
LOCUS
DEFINITION Sequence 221 from Patent WO0179548.
AX288459
ACCESSION AX288459
VERSION AX288459.1 GI:17050142
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct
artificial sequences.
REFERENCE 1
AUTHORS Barany, F., Zirvi, M., Gerry, N.P., Favis, R. and Kliman, R.
TITLE Method of designing addressable array for detection of nucleic acid
sequence differences using ligase detection reaction
JOURNAL Patent: WO 0179548-A 221 25-OCT-2001;
CORNELL RESEARCH FOUNDATION, INC. (US)
FEATURES source Location/Qualifiers
1..24 /organism="synthetic construct"
/db_xref="taxon:32630"
/note="Hypothetical Probe Sequence"
BASE COUNT 1 a 7 c 7 g 9 t
ORIGIN
Query Match 65.6%; Score 11.8; DB 6; Length 24;
Best Local Similarity 60.0%; Pred. No. 9.2e+04;
Matches 9; Conservative 4; Mismatches 2; Indels 0; Gaps 0;
Qy 3 UGGCCUAUCGGUGCG 17
Db 6 TGCCTATCTGTGGC 20
RESULT 13
AX447444
LOCUS
DEFINITION Sequence 3899 from Patent WO0216649.
AX447444
ACCESSION AX447444
VERSION AX447444.1 GI:21696343
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct
artificial sequences.
REFERENCE 1
AUTHORS Gunderson, K.
TITLE Probes and decoder oligonucleotides
JOURNAL Patent: WO 0216649-A 3899 28-FEB-2002;
Illumina, Inc. (US)
FEATURES source Location/Qualifiers
1..24 /organism="synthetic construct"
/db_xref="taxon:32630"
/note="Computer Generated Probe Sequence."
BASE COUNT 6 a 6 c 8 g 4 t
ORIGIN
Query Match 65.6%; Score 11.8; DB 6; Length 24;
Best Local Similarity 73.3%; Pred. No. 9.2e+04;
Matches 11; Conservative 2; Mismatches 2; Indels 0; Gaps 0;
Qy 4 GGCCUAUCGGUGCGA 18
Db 2 GGCCTAGAGTGGCA 16
RESULT 14
AX29209
LOCUS
DEFINITION DNA probe from patent WO9111459.
AX29209
ACCESSION AX29209
VERSION AX29209.1 GI:1248930
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct
artificial sequences.
REFERENCE 1
AUTHORS
TITLE
JOURNAL
FEATURES source Location/Qualifiers
1..30 /organism="synthetic construct"
/db_xref="taxon:32630"
BASE COUNT 5 a 6 c 13 g 6 t
ORIGIN

ORIGIN

Query Match 65.6%; Score 11.8; DB 6; Length 30;
 Best Local Similarity 73.3%; Pred. NO. 9.2e+04;
 Matches 11; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 2 AUGGCCUAUCGGUGC 16
 | | | | | | | | | |
 Db 1 ACGGCTAGCGGTGC 15

RESULT 15

A29212
 LOCUS 30 bp DNA linear PAT 30-JUN-1995
 DEFINITION Oligonucleotide OAB1088 from patent WO9111459.

ACCESSION A29212
 VERSION A29212.1 GI:1248933

KEYWORDS
 SOURCE synthetic construct.
 ORGANISM synthetic construct
 artificial sequences.

FEATURES
 source Location/Qualifiers
 1..30
 /organism="synthetic construct"
 /db_xref="taxon:32630"
 BASE COUNT 5 a 6 c 13 g 6 t
 ORIGIN

Query Match 65.6%; Score 11.8; DB 6; Length 30;
 Best Local Similarity 73.3%; Pred. NO. 9.2e+04;
 Matches 11; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 2 AUGGCCUAUCGGUGC 16
 | | | | | | | | | |
 Db 1 ACGGCTAGCGGTGC 15

Search completed: July 6, 2003, 14:51:13
 Job time : 593.818 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: July 6, 2003, 14:25:15 ; Search time 193.091 Seconds
(without alignments)
209.932 Million cell updates/sec

Title: US-09-780-929-98
Perfect score: 18
Sequence: 1 aauGCCuauCGGUGCGA 18

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 2185239 seqs, 1125999159 residues

Total number of hits satisfying chosen parameters: 2274872

Minimum DB seq length: 0
Maximum DB seq length: 60

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : N_Geneseq_101002.*
1: /SID22/gcgdata/geneseq/geneseq-emb1/NA1980.DAT.*
2: /SID22/gcgdata/geneseq/geneseq-emb1/NA1981.DAT.*
3: /SID22/gcgdata/geneseq/geneseq-emb1/NA1982.DAT.*
4: /SID22/gcgdata/geneseq/geneseq-emb1/NA1983.DAT.*
5: /SID22/gcgdata/geneseq/geneseq-emb1/NA1984.DAT.*
6: /SID22/gcgdata/geneseq/geneseq-emb1/NA1985.DAT.*
7: /SID22/gcgdata/geneseq/geneseq-emb1/NA1986.DAT.*
8: /SID22/gcgdata/geneseq/geneseq-emb1/NA1987.DAT.*
9: /SID22/gcgdata/geneseq/geneseq-emb1/NA1988.DAT.*
10: /SID22/gcgdata/geneseq/geneseq-emb1/NA1989.DAT.*
11: /SID22/gcgdata/geneseq/geneseq-emb1/NA1990.DAT.*
12: /SID22/gcgdata/geneseq/geneseq-emb1/NA1991.DAT.*
13: /SID22/gcgdata/geneseq/geneseq-emb1/NA1992.DAT.*
14: /SID22/gcgdata/geneseq/geneseq-emb1/NA1993.DAT.*
15: /SID22/gcgdata/geneseq/geneseq-emb1/NA1994.DAT.*
16: /SID22/gcgdata/geneseq/geneseq-emb1/NA1995.DAT.*
17: /SID22/gcgdata/geneseq/geneseq-emb1/NA1996.DAT.*
18: /SID22/gcgdata/geneseq/geneseq-emb1/NA1997.DAT.*
19: /SID22/gcgdata/geneseq/geneseq-emb1/NA1998.DAT.*
20: /SID22/gcgdata/geneseq/geneseq-emb1/NA1999.DAT.*
21: /SID22/gcgdata/geneseq/geneseq-emb1/NA2000.DAT.*
22: /SID22/gcgdata/geneseq/geneseq-emb1/NA2001A.DAT.*
23: /SID22/gcgdata/geneseq/geneseq-emb1/NA2001B.DAT.*
24: /SID22/gcgdata/geneseq/geneseq-emb1/NA2002.DAT.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	18	100.0	18	22 AAS12348	DNA encoding deoxy
2	18	100.0	29	22 AAS12381	Class IV ribozyme.
C 3	13.8	76.7	30	24 AAS19241	CMV mutagenic adap
4	13.4	74.4	60	24 ABN47743	Human spliced tran
5	12.8	71.1	20	24 ABI95945	Capture oligonucle
6	12.8	71.1	24	24 ABI88460	Capture oligonucle
C 7	12.8	71.1	24	24 ABI88461	Capture oligonucle
8	12.4	68.9	41	22 AAC87378	Staphylococcus aur
9	12.2	67.8	20	24 ABI93941	Capture oligonucle

C 10	12.2	67.8	24	22 AAS08711	Human PD-ATP-bindin
C 11	12.2	67.8	24	24 ABI84452	Capture oligonucle
C 12	12.2	67.8	24	24 ABI84453	Capture oligonucle
C 13	12.2	67.8	29	20 AAX98255	PCR primer used to
C 14	12.2	67.8	40	17 AAT70787	Stenotic carotid a
C 15	12.2	67.8	53	16 AAT21824	Human gene signatu
C 16	12.2	67.8	60	24 ABN32869	Human spliced tran
C 17	12	66.7	22	24 ABA04884	TT virus PCR prime
C 18	11.8	65.6	20	20 AAX94010	PCR primer used to
C 19	11.8	65.6	20	21 AAZ44197	Murine cerebral ne
C 20	11.8	65.6	20	24 ABI93134	Capture oligonucle
C 21	11.8	65.6	22	20 AAZ07252	Wild-type oligo h1
C 22	11.8	65.6	22	20 AAZ07306	Human telomerase R
C 23	11.8	65.6	24	24 ABQ03892	Oligonucleotide ad
C 24	11.8	65.6	24	24 ABI82838	Capture oligonucle
C 25	11.8	65.6	24	24 ABI82839	Capture oligonucle
C 26	11.8	65.6	24	24 ABI92104	Capture oligonucle
C 27	11.8	65.6	24	24 ABI92105	Capture oligonucle
C 28	11.8	65.6	30	12 AAQ13310	Probe OAB 1088 for
C 29	11.8	65.6	39	22 AAI68437	P.carinii 18S rRNA
C 30	11.8	65.6	40	21 AAA62656	Beta-lactamase gen
C 31	11.6	64.4	22	22 AAD11273	Mycobacterium 16S
C 32	11.6	64.4	24	22 AAD11272	Mycobacterium 16S
C 33	11.6	64.4	25	22 AAD11271	Mycobacterium 16S
C 34	11.6	64.4	27	19 RAV55972	Human cytokine rec
C 35	11.6	64.4	29	16 AAQ99711	Primer corresp. to
C 36	11.6	64.4	29	17 AAT10032	MDGF antisense pri
C 37	11.6	64.4	30	16 AAT04920	Mammalian stem cel
C 38	11.6	64.4	30	16 AAO99710	Primer for reverse
C 39	11.6	64.4	30	17 AAT10031	MDGF cDNA/antisens
C 40	11.6	64.4	30	20 AAX36536	PCR primer for hum
C 41	11.6	64.4	30	21 AAX13756	Stem cell factor u
C 42	11.6	64.4	30	21 AAZ55799	Reverse transcript
C 43	11.6	64.4	30	22 AAS10451	Human stem cell fa
C 44	11.6	64.4	30	22 AAH41335	Universal stem cel
C 45	11.6	64.4	30	22 AAS04115	Human SCF (stem ce

ALIGNMENTS

RESULT 1

AAS12348
ID AAS12348 standard; DNA; 18 BP.

XX AAS12348;

XX 21-NOV-2001 (first entry)

XX DNA encoding deoxyribozyme #8.

XX Deoxyribozyme; cytosstatic; endonuclease; RNA cleavage; DNA cleavage;
XX gene therapy; plant; fungus; bacteria; mammal; ribozyme; ss.

XX Synthetic.

XX WO200159102-A2.

XX 16-AUG-2001.

XX 08-FEB-2001; 2001WO-US04223.

XX 08-FEB-2000; 2000US-0181360.

XX 31-MAR-2000; 2000US-0193646.

XX (RIBO-) RIBOZYME PHARM INC.

XX (UYIA) UNIV YALE.

XX Breaker R, Belgelman L, Emilsson G;

XX WPI; 2001-536526/59.

XX New nucleic acids with endonuclease activity, such as ribozymes and

PT nucleozymes, for modulating gene expression in a plant, mammalian,
PT bacterial or fungal cell -
PS Claim 49; Page 77; 96pp; English.
XX
CC The invention relates to nucleic acid molecules with endonuclease
CC activity, which are particularly useful for cleavage of RNA or DNA.
CC The nucleic acids are used in a pharmaceutical composition and are used
CC to modulate expression of a gene in a plant, mammalian, bacterial or
CC fungal cell. They are used to cleave a separate nucleic acid, preferably
CC RNA. The nucleic acids are used to inhibit gene expression and/or cell
CC proliferation, and can be used to treat a disease or condition. More
CC than one nucleic acid can be independently targeted to the same or
CC different sites in a cell. The nucleic acids may be used to study DNA.
CC The modifications to the nucleic acids optimises their catalytic activity
CC and can maintain or enhance their activity. They exhibit a high degree
CC of specificity for RNA. The present sequence represents the coding
CC sequence of deoxyribozyme #8 used in the method of the invention.
XX
XX Sequence 18 BP; 4 A; 4 C; 6 G; 4 U; 0 other;
SQ
Query Match 100.0%; Score 18; DB 22; Length 18;
Best Local Similarity 100.0%; Pred. No. 0.63;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 AAUGGCCUAUCGGUGCGA 18
DB 1 AAUGGCCUAUCGGUGCGA 18
RESULT 2
AAS12381
ID AAS12381 standard; RNA; 29 BP.
XX
AC AAS12381;
XX
XX 21-NOV-2001 (first entry)
XX
XX Class IV ribozyme.
XX
XX Deoxyribozyme; cytotstatic; endonuclease; RNA cleavage; DNA cleavage;
XX gene therapy; plant; fungus; bacteria; mammal; ribozyme; ss.
XX Synthetic.
XX
XX Key Location/Qualifiers
XX misc_binding 1..8
XX /*tag= a
XX /*note= "Forms double-stranded region with bases 15
XX to 8 of AAS12374"
XX
XX misc_binding 25..29
XX /*tag= b
XX /*note= "Forms double-stranded region with bases 5
XX to 1 of AAS12374"
XX
XX WO200159102-A2.
XX
XX 16-AUG-2001.
XX
XX 08-FEB-2001; 2001WO-US04223.
XX
XX 08-FEB-2000; 2000US-0181360.
XX
XX 31-MAR-2000; 2000US-0193646.
XX
XX (RIBO-) RIBOZYME PHARM INC.
XX PA (UYVA) UNIV YALE.
XX
XX Breaker R, Beigelman L, Emilsson G;
XX WPI; 2001-536526/59.
XX
XX New nucleic acids with endonuclease activity, such as ribozymes and
PT nucleozymes, for modulating gene expression in a plant, mammalian,
PT

PT bacterial or fungal cell -
XX
XX Example 1; Fig 9; 96pp; English.
XX
CC The invention relates to nucleic acid molecules with endonuclease
CC activity, which are particularly useful for cleavage of RNA or DNA.
CC The nucleic acids are used in a pharmaceutical composition and are used
CC to modulate expression of a gene in a plant, mammalian, bacterial or
CC fungal cell. They are used to cleave a separate nucleic acid, preferably
CC RNA. The nucleic acids are used to inhibit gene expression and/or cell
CC proliferation, and can be used to treat a disease or condition. More
CC than one nucleic acid can be independently targeted to the same or
CC different sites in a cell. The nucleic acids may be used to study DNA.
CC The modifications to the nucleic acids optimises their catalytic activity
CC and can maintain or enhance their activity. They exhibit a high degree
CC of specificity for RNA. The present sequence represents the Class IV
CC ribozyme, used in an example which demonstrates the method of
XX the invention.
XX
XX Sequence 29 BP; 6 A; 7 C; 11 G; 5 U; 0 other;
SQ
Query Match 100.0%; Score 18; DB 22; Length 29;
Best Local Similarity 100.0%; Pred. No. 0.67;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 AAUGGCCUAUCGGUGCGA 18
DB 8 AAUGGCCUAUCGGUGCGA 25
RESULT 3
AAS19241/c
ID AAS19241 standard; DNA; 30 BP.
XX
AC AAS19241;
XX
XX 26-MAR-2002 (first entry)
XX
XX CMV mutagenic adaptor.
XX
XX T0; ds; terminator; pGA; DNA vaccine; anti-HIV; virucide;
XX Human Immunodeficiency Virus; HIV; Gag; HIV gp120; HIV Pol; HIV Env;
XX HIV VLP; measles fusion protein; measles haemagglutinin; CMV; adaptor;
XX measles nucleoprotein; influenza haemagglutinin; C3d gene;
XX cell-mediated immune response; humoral immune response; infection.
XX
XX Human cytomegalovirus.
XX
XX WO200192470-A2.
XX
XX 06-DEC-2001.
XX
XX 02-MAR-2001; 2001WO-US06795.
XX
XX 02-MAR-2000; 2000US-186364P.
XX
XX 01-DEC-2000; 2000US-251083P.
XX
XX (UYEM-) UNIV EMORY.
XX
XX Robinson HL, Smith JM, Ross TM, Bright RA, Hua J, Ellenberger D;
XX WPI; 2002-075465/10.
XX
XX Novel pGA vector useful for immunising patient against measles,
XX influenza has termination sequence encoding lambda T0 terminator and a
XX eukaryotic transcription cassette with vaccine insert encoding
XX immunogens of pathogens -
XX
XX Example 2; Page 43; 174pp; English.
XX
XX The invention relates to a vector (a pGA construct) comprising a
XX termination sequence coding for the lambda T0 terminator, a prokaryotic
XX origin of replication, a selectable marker gene and a eukaryotic

transcription cassette comprising a vaccine insert encoding one or more immunogens derived from a pathogen e.g. Human Immunodeficiency Virus (HIV) Gag, HIV gp120, HIV Pol, HIV Env, HIV VLP, or its mutants, measles fusion protein, measles haemagglutinin, measles nucleoprotein, influenza haemagglutinin, or its mutants, or subsequences, and optionally at least one C3d gene, is useful for immunising or treating a patient, when administered by an intramuscular or intradermal route. The immunisation methods using pGA elicit both cell-mediated and humoral immune responses that may limit the infection, spread or growth of the pathogen and result in protection against subsequent challenge against the pathogen. The terminator sequence present prevents read-through from the kanamycin cassette into vaccine sequences while the plasmid is being produced in bacteria. Prevention of transcriptional read-through stabilises vaccine insert sequences by limiting the exposure of secondary structures that can be recognised by bacterial endonucleases. The present sequence is an adaptor for introducing a ClaI site into the cytomegalovirus (CMV) promoter of pGA2.

XX SQ Sequence 30 BP; 7 A; 12 C; 6 G; 5 T; 0 other;

Query Match 76.7%; Score 13.8; DB 24; Length 30;
Best Local Similarity 64.7%; Pred. No. 1.9e+02; Indels 0; Gaps 0;
Matches 11; Conservative 4; Mismatches 2; Indels 0; Gaps 0;

QY 2 AUGGCCUAUCGGUGCGA 18
I:III:IIII:IIII
Db 25 ATGGCGTATCGATCGGA 9

RESULT 4

ABN47743
ID ABN47743 standard; DNA; 60 BP.

XX AC ABN47743;

XX DT 15-JUL-2002 (first entry)

XX DE Human spliced transcript detection oligonucleotide SEQ ID NO:20491.

XX KW Human; mouse; rat; splice transcript; detection; RNA transcript;
splice variant; transcriptome; oligonucleotide library; ss.

XX OS Homo sapiens.

XX PN WO200210449-A2.

XX PD 07-FEB-2002.

XX PF 20-JUL-2001; 2001WO-IB01903.

XX PR 28-JUL-2000; 2000US-221607P.

XX PR 02-MAY-2001; 2001US-287724P.

XX PA (COMP-) COMPUGEN INC.

XX PI Shoshan A, Wasserman A, Mintz E, Mintz L, Faigler S;

XX DR WPI; 2002-257383/30.

XX PT New oligonucleotide libraries comprising oligonucleotides which selectively hybridize to mRNAs transcribed from a transcription unit of a genome, useful for detecting tissue-, pathology-, and developmental-specific genes -

XX PS Example 1; SEQ ID 20491; 47pp; English.

XX CC The present invention describes oligonucleotide libraries for detecting messenger RNAs that populate a (sub-)transcriptome, where the (sub-)transcriptome comprises messenger RNAs transcribed from multiple transcription units that populate a genome. The library comprises several oligonucleotides, each capable of hybridising selectively to a set of messenger RNAs transcribed from a given transcription unit of the genome, which encodes one or more messenger RNA splice variants.

CC The oligonucleotide libraries are useful for detecting mRNAs from a biological sample, in expression profiling studies, in qualitatively or quantitatively characterising the corresponding transcriptome, and in detecting RNA transcripts and splice variants of human or animal transcriptomes. The libraries may also be used as specialised mini libraries to detect transcripts of a sub-transcriptome under a particular biological or pathological state, and so allowing the detection of tissue- and pathology-specific genes such as those genes only expressed in specific tissue under a specific pathological condition; to detect developmental specific genes; and to detect RNA transcripts and splice variants of a transcriptome of a patient suffering from a particular disorder. ABN27253 to ABN59589 represent oligonucleotide sequences from rats, humans and mice, which are used in the exemplification of the present invention.

CC N.B. The sequence data for this patent did not form part of the printed specification, but was obtained in electronic format directly from WIPO at ftp.wipo.int/pub/published_pct_sequences.

XX SQ Sequence 60 BP; 16 A; 18 C; 13 G; 13 T; 0 other;

Query Match 74.4%; Score 13.4; DB 24; Length 60;
Best Local Similarity 66.7%; Pred. No. 3.6e+02; Indels 0; Gaps 0;
Matches 10; Conservative 4; Mismatches 1; Indels 0; Gaps 0;

QY 1 AAUGGCCUAUCGGUG 15
I:IIII:IIII:IIII
Db 1 ATGGCCTATCGGTG 15

RESULT 5

ABI95945

ID ABI95945 standard; DNA; 20 BP.

XX AC ABI95945;

XX DT 16-FEB-2002 (first entry)

XX DE Capture oligonucleotide Zip ID#3032 oligo #9.

XX KW Human; K-ras; PCR primer; probe; capture probe; mutation detection;
ligase detection reaction; LDR; p53; BRCA1; BRCA2; infectious disease;
infection; 21 hydroxylase deficiency; Turner Syndrome; obesity;
cancer; oncogene; tumour suppressor; human papillomavirus; forensic;
environmental monitoring; food industry; feed industry; ss.

XX OS Synthetic.

XX PN WO200179548-A2.

XX PD 25-OCT-2001.

XX PF 04-APR-2001; 2001WO-US10958.

XX PR 14-APR-2000; 2000US-197271P.

XX PA (CORR) CORNELL RES FOUND INC.

XX PI Barany F, Zirvi M, Gerry NP, Favis R, Kilman R;

XX DR WPI; 2002-034366/04.

XX PT Designing capture oligonucleotide probes for use on a support to which complementary oligonucleotides hybridize with little mismatch -

XX PS Example 5; Fig 29; 300pp; English.

XX CC The present invention describes a method (M1) for designing capture oligonucleotide probes (I) for use on a support to which complementary oligonucleotide probes (II) will hybridise with little mismatch, where (I) have melting temperatures within a narrow range. The method is useful for detecting infectious diseases caused by bacterial infectious agents e.g. Salmonella, Listeria monocytogenes and Haemophilus influenzae, fungal infectious agents e.g. Cryptococcus neoformans, Candida albicans and

CC Aspergillus fumigatus, viruses e.g. T-cell lymphocytotropic virus,
 CC Epstein-Barr virus and polio virus, and parasitic infectious agents
 CC selected from Onchocerca volvulus, Entamoeba histolytica and Dracunculus
 CC medinensis. The method is also useful for detecting genetic diseases such
 CC as 21 hydroxylase deficiency, Turner Syndrome and obesity defects.
 CC Detecting cancer involving oncogenes, tumour suppressor genes, or genes
 CC involved in DNA amplification, replication, recombination or repair, the
 CC cancer is specifically associated with a gene selected from BRCA1 gene,
 CC p53 gene, human papillomavirus types 16 and 18 and liver cancers. The
 CC method is also used for environmental monitoring, forensics and the food
 CC and feed industry, detecting comprises scanning (using e.g. a scanning
 CC electron microscope and infrared microscope) the support at the
 CC particular sites and identifying if ligation of the oligonucleotide probe
 CC sets occurred and correlating (using a computer) identified ligation to a
 CC presence or absence of the target nucleotide sequences. AB182074 to
 CC AB197546 represent oligonucleotide sequences used in the exemplification
 CC of the present invention.

XX SQ Sequence 20 BP; 2 A; 6 C; 6 G; 6 T; 0 other;

Query Match 71.1%; Score 12.8; DB 24; Length 20;
 Best Local Similarity 68.8%; Pred. No. 7.2e+02;
 Matches 11; Conservative 3; Mismatches 2; Indels 0; Gaps 0;

Qy 2 AUGGCCUAUCGGUGCG 17
 1 111 :1:111:111
 Db 2 ACGGCTTATCGGTGCG 17

RESULT 6

AB188460
 ID AB188460 standard; DNA; 24 BP.

AC AB188460;

XX 15-FEB-2002 (first entry)

DE Capture oligonucleotide Zip ID#3032 oligo #1.

XX Human; K-ras; PCR primer; probe; capture probe; mutation detection;
 KW ligase detection reaction; LDR; p53; BRCA1; BRCA2; infectious disease;
 KW infection; 21 hydroxylase deficiency; Turner Syndrome; obesity;
 KW cancer; oncogene; tumour suppressor; human papillomavirus; forensic;
 KW environmental monitoring; food industry; feed industry; ss.

OS Synthetic.

XX WO200179548-A2.

XX 25-OCT-2001.

XX 04-APR-2001; 2001WO-US10958.

XX 14-APR-2000; 2000US-197271P.

XX (CORR) CORNELL RES FOUND INC.

XX Barany F, Zirvi M, Gerry NP, Favis R, Kliman R;

XX WPI; 2002-034366/04.

PT Designing capture oligonucleotide probes for use on a support to which
 PT complementary oligonucleotides hybridize with little mismatch -

XX Example 5; Fig 25; 300pp; English.

XX The present invention describes a method (M1) for designing capture
 CC oligonucleotide probes (I) for use on a support to which complementary
 CC oligonucleotide probes (II) will hybridize with little mismatch, where
 CC (I) have melting temperatures within a narrow range. The method is useful
 CC for detecting infectious diseases caused by bacterial infectious agents
 CC e.g. Salmonella, Listeria monocytogenes and Haemophilus influenza, fungal
 CC infectious agents e.g. Cryptococcus neoformans, Candida albicans and

CC Aspergillus fumigatus, viruses e.g. T-cell lymphocytotropic virus,
 CC Epstein-Barr virus and polio virus, and parasitic infectious agents
 CC selected from Onchocerca volvulus, Entamoeba histolytica and Dracunculus
 CC medinensis. The method is also useful for detecting genetic diseases such
 CC as 21 hydroxylase deficiency, Turner Syndrome and obesity defects.
 CC Detecting cancer involving oncogenes, tumour suppressor genes, or genes
 CC involved in DNA amplification, replication, recombination or repair, the
 CC cancer is specifically associated with a gene selected from BRCA1 gene,
 CC p53 gene, human papillomavirus types 16 and 18 and liver cancers. The
 CC method is also used for environmental monitoring, forensics and the food
 CC and feed industry, detecting comprises scanning (using e.g. a scanning
 CC electron microscope and infrared microscope) the support at the
 CC particular sites and identifying if ligation of the oligonucleotide probe
 CC sets occurred and correlating (using a computer) identified ligation to a
 CC presence or absence of the target nucleotide sequences. AB182074 to
 CC AB197546 represent oligonucleotide sequences used in the exemplification
 CC of the present invention.

XX SQ Sequence 24 BP; 2 A; 7 C; 7 G; 8 T; 0 other;

Query Match 71.1%; Score 12.8; DB 24;
 Best Local Similarity 68.8%; Pred. No. 7.3e+02;
 Matches 11; Conservative 3; Mismatches 2; Indels 0; Gaps 0;

Qy 2 AUGGCCUAUCGGUGCG 17
 1 111 :1:111:111
 Db 2 ACGGCTTATCGGTGCG 17

RESULT 7

AB188461/G

ID AB188461 standard; DNA; 24 BP.

AC AB188461;

XX 15-FEB-2002 (first entry)

DE Capture oligonucleotide Zip ID#3032 oligo #2.

XX Human; K-ras; PCR primer; probe; capture probe; mutation detection;
 KW ligase detection reaction; LDR; p53; BRCA1; BRCA2; infectious disease;
 KW infection; 21 hydroxylase deficiency; Turner Syndrome; obesity;
 KW cancer; oncogene; tumour suppressor; human papillomavirus; forensic;
 KW environmental monitoring; food industry; feed industry; ss.

OS Synthetic.

XX WO200179548-A2.

XX 25-OCT-2001.

XX 04-APR-2001; 2001WO-US10958.

XX 14-APR-2000; 2000US-197271P.

XX (CORR) CORNELL RES FOUND INC.

XX Barany F, Zirvi M, Gerry NP, Favis R, Kliman R;

XX WPI; 2002-034366/04.

PT Designing capture oligonucleotide probes for use on a support to which
 PT complementary oligonucleotides hybridize with little mismatch -

XX Example 5; Fig 25; 300pp; English.

XX The present invention describes a method (M1) for designing capture
 CC oligonucleotide probes (I) for use on a support to which complementary
 CC oligonucleotide probes (II) will hybridize with little mismatch, where
 CC (I) have melting temperatures within a narrow range. The method is useful
 CC for detecting infectious diseases caused by bacterial infectious agents
 CC e.g. Salmonella, Listeria monocytogenes and Haemophilus influenza, fungal
 CC infectious agents e.g. Cryptococcus neoformans, Candida albicans and

Aspergillus fumigatus, viruses e.g. T-cell lymphocytotropic virus, Epstein-Barr virus and polio virus, and parasitic infectious agents selected from Onchocerca volvulus, Entamoeba histolytica and Dracunculus medinensis. The method is also useful for detecting genetic diseases such as 21 hydroxylase deficiency, Turner Syndrome and obesity defects. Detecting cancer involving oncogenes, tumour suppressor genes, or genes involved in DNA amplification, replication, recombination or repair, the cancer is specifically associated with a gene selected from BRCA1 gene, p53 gene, human papillomavirus types 16 and 18 and liver cancers. The method is also used for environmental monitoring, forensics and the food and feed industry, detecting comprises scanning (using e.g. a scanning electron microscope and infrared microscope) the support at the particular sites and identifying if ligation of the oligonucleotide probe sets occurred and correlating (using a computer) identified ligation to a presence or absence of the target nucleotide sequences. ABI82074 to ABI97546 represent oligonucleotide sequences used in the exemplification of the present invention.

Sequence 24 BP; 8 A; 7 C; 7 G; 2 T; 0 other;

Query Match 71.1%; Score 12.8; DB 24; Length 24;

Best Local Similarity 68.8%; Pred. No. 7.3e+02;

Matches 11; Conservative 3; Mismatches 2; Indels 0; Gaps 0;

Qy 2 AUGCCUAUCGGUGCG 17

Db 23 ACGGTTATCGGTGCG 8

RESULT 8

AAC87378

ID AAC87378 standard; DNA; 41 BP.

XX AAC87378;

AC AAC87378;

DT 09-MAR-2001 (first entry)

XX Staphylococcus aureus Spa domain D antisense PCR primer, PC3H AS.

DE Staphylococcus aureus

XX Spa domain D; randomised library; VH3 Ig-Fab fragment; immunoglobulin;

KW Spa mutant; superantigen; altered specificity; apoptosis inducer;

KW energy inducer; B-lymphocyte subset; B-cell; lymphoma; leukaemia;

KW autoimmune disease; idiopathic thrombocytopenia; rheumatoid arthritis;

KW systemic lupus erythematosus; SLE; autoimmune thyroiditis; diabetes.

KW antibody purification; PCR primer; ss.

XX Staphylococcus aureus.

OS Staphylococcus aureus.

XX WO200069457-A1.

PN 23-NOV-2000.

XX 15-MAY-2000; 2000WO-US13402.

PF 15-MAY-1999; 99US-0134386.

XX (UYCA-) UNIV CALIFORNIA SAN DIEGO.

XX Silverman GJ;

XX WPI; 2001-031886/04.

XX New staphylococcal protein A variant, useful for treating diabetes and

PT rheumatoid arthritis, exhibits binding specificity for

PT immunoglobulin-Fab domain and comprises variations in amino acid

XX sequence of staphylococcal protein A domain D

XX Example 12; Page 58; 88pp; English.

XX The invention relates to staphylococcal protein A (SpA) variants which

CC exhibit altered binding specificity for an immunoglobulin Fab (Ig-Fab)

CC fragment relative to native SpA. The SpA variants of the invention have

CC one or more amino acid substitution in the SpA VH3 Ig-Fab binding region

XX

(i.e., SpA domain D) relative to the native SpA. The SpA variants are inducers of autoreactive B cell, leukaemic or lymphoma cell apoptosis or energy. The SpA variants of the invention are useful for detecting the presence of a certain Ig-Fab-expressing lymphocyte subset in a sample of lymphocytes. The SpA variants may therefore be used in the diagnosis of some forms of leukaemia. The SpA variants may also be administered to an individual with an abnormally high number of a certain lymphocyte subset to reduce the number of that lymphocyte subset. The SpA variants are also useful for purifying monoclonal or polyclonal antibodies from serum, plasma, tissue culture or other sources. The SpA variants that exhibit enhanced clonogenicity or other properties in vitro or in vivo, such as the ability to delete undesirable neoplastic B-cells or pathogenic B-cells that are responsible for the production of disease-causing autoantibodies, are useful as therapeutic agents. Therapeutic SpA variants that can bind to Fab on the B-cell receptor of an autoreactive B-cell or leukaemic or lymphoma cell can induce energy, apoptosis, or deletion by other mechanisms. By engineering the interaction between SpA variants and Ig-Fab according to the invention, variant SpA with specially tailored Fab-binding specificities can be selected that target pathogenic neoplastic B cell populations or autoreactive B-cell clones. The SpA variants are therefore useful for treating conditions that are linked to disease-associated B-cells such as idiopathic thrombocytopenia, rheumatoid arthritis, systemic lupus erythematosus (SLE), autoimmune thyroiditis, or diabetes. Engineered B cell superantigens (such as the SpA variants of the invention) bind to immunoglobulin receptors on B-cells in a manner that is distinct from antigen binding by antibodies. Therefore, the SpA variants which have superantigen properties can target much larger populations of Ig-expressing B cells in vivo compared with targeting using a specific antigen. Sequences AAC87372-C87378 represent nucleic acid sequences used in the construction of a randomised SpA domain D library in an exemplification of the invention. Sequences AAC87375-C87378 represent SpA domain D PCR primers used in library construction.

Sequence 41 BP; 5 A; 9 C; 19 G; 8 T; 0 other;

Query Match 68.9%; Score 12.4; DB 22; Length 41;

Best Local Similarity 64.3%; Pred. No. 1.3e+03;

Matches 9; Conservative 4; Mismatches 1; Indels 0; Gaps 0;

Qy 3 UGGCCUAUCGGUGCG 16

Db 23 TGGCTTTCGGTGC 36

RESULT 9

ABI93941

ID ABI93941 standard; DNA; 20 BP.

XX ABI93941;

AC ABI93941;

XX 16-FEB-2002 (first entry)

XX Capture oligonucleotide Zip ID#1028 oligo #9.

DE Human; K-ras; PCR primer; probe; capture probe; mutation detection;

XX ligase detection reaction; LDR; p53; BRCA1; BRCA2; infectious disease;

KW infection; 21 hydroxylase deficiency; Turner Syndrome; obesity;

KW cancer; oncogene; tumour suppressor; human papillomavirus; forensic;

KW environmental monitoring; food industry; feed industry; ss.

XX Synthetic.

OS WO200179548-A2.

PN 25-OCT-2001.

XX 04-APR-2001; 2001WO-US10958.

XX 14-APR-2000; 2000US-197271P.

XX (CORR) CORNELL RES FOUND INC.

PA

XX

```
PI Barany F, Zirvi M, Gerry NP, Favis R, Kliman R;
XX WPI; 2002-034366/04.
XX
XX Designing capture oligonucleotide probes for use on a support to which
XX complementary oligonucleotides hybridize with little mismatch -
XX
XX Example 5; Fig 29; 300pp; English.
XX
XX The present invention describes a method (M1) for designing capture
XX oligonucleotide probes (I) for use on a support to which complementary
XX oligonucleotide probes (II) will hybridize with little mismatch, where
XX (I) have melting temperatures within a narrow range. The method is useful
XX for detecting infectious diseases caused by bacterial infectious agents
XX e.g. Salmonella, Listeria monocytogenes and Haemophilus influenzae, fungal
XX infectious agents e.g. Cryptococcus neoformans, Candida albicans and
XX Aspergillus fumigatus, viruses e.g. T-cell lymphocytotropic virus,
XX Epstein-Barr virus and polio virus, and parasitic infectious agents
XX selected from Onchocerca volvulus, Entamoeba histolytica and Dracunculus
XX medinensis. The method is also useful for detecting genetic diseases such
XX as 21 hydroxylase deficiency, Turner Syndrome and obesity defects.
XX Detecting cancer involving oncogenes, tumour suppressor genes, or genes
XX involved in DNA amplification, replication, recombination or repair, the
XX cancer is specifically associated with a gene selected from BRCA1 gene,
XX p53 gene, human papillomavirus types 16 and 18 and liver cancers. The
XX method is also used for environmental monitoring, forensics and the food
XX and feed industry, detecting comprises scanning (using e.g. a scanning
XX electron microscope and infrared microscope) the support at the
XX particular sites and identifying if ligation of the oligonucleotide probe
XX sets occurred and correlating (using a computer) identified ligation to a
XX presence or absence of the target nucleotide sequences. ABI82074 to
XX ABI97546 represent oligonucleotide sequences used in the exemplification
XX of the present invention.
XX
XX Sequence 20 BP; 7 A; 6 C; 4 G; 3 T; 0 other;
SQ Query Match 67.8%; Score 12.2; DB 24; Length 20;
Best Local Similarity 64.7%; Pred. No. 1.6e+03;
Matches 11; Conservative 3; Mismatches 3; Indels 0; Gaps 0;

QY 2 AUGGCCUAUCGGUGCGA 18
I : I I I I I I I I I I
D b 3 ATGACCAATCGATCGCA 19

RESULT 10
AAS08711/c
ID AAS08711 standard; DNA; 24 BP.
AC AAS08711;
XX
XX 26-SEP-2001 (first entry)
XX
XX Human PD-ATP-binding cassette (PD-ABC) cDNA reverse PCR primer #2.
XX
XX PD-ATP-binding cassette; PD-ABC; chromosome 19p13.3; spleen; thymus; ss;
XX peripheral blood leukocyte; bone marrow; lymph node; dyslipidaemia;
XX cardiovascular disorder; inflammatory disorder; abnormal calcium flux;
XX epilepsy; coronary artery disease; Tangier's disease; atherosclerosis;
XX familial high-density lipoprotein deficiency; fatty liver disease;
XX atherosclerosis; diabetes; insulin resistance; obesity; drug screening;
XX alcoholism; retinal degeneration; hypertension; vascular disease;
XX PCR primer.
XX
XX Synthetic.
XX
XX WO200153490-A1.
XX
XX 26-JUL-2001.
XX
XX 23-JAN-2001; 2001WO-US02191.
XX
XX 24-JAN-2000; 2000US-0177899.
XX
XX
```

```
PR 30-JUN-2000; 2000US-0215405.
XX (WARN ) WARNER LAMBERT CO.
XX
XX Johns MA, Tafuri SR, Wang M;
XX WPI; 2001-442259/47.
XX
XX New Human PD-ABC DNA molecules and proteins for diagnosis and treatment
XX of dyslipidaemia, epilepsy and diseases related to abnormal calcium flux
XX
XX Disclosure; Page 34; 77pp; English.
XX
XX The sequence represents a PCR primer used for isolation of cDNA encoding
XX human PD-ATP-binding cassette (PD-ABC) protein. PD-ABC maps to chromosome
XX 19p13.3 and is expressed in various tissues including spleen, thymus,
XX peripheral blood leukocytes, bone marrow and lymph nodes. The PD-ABC DNA
XX molecules and proteins are used to diagnose and treat cardiovascular
XX disorders, inflammatory disorders, dyslipidaemia, epilepsy, diseases
XX related to abnormal calcium flux, coronary artery disease, Tangier's
XX disease, familial high-density lipoprotein deficiency, atherosclerosis,
XX diabetes, fatty liver disease, insulin resistance, obesity, alcoholism,
XX retinal degeneration, hypertension and vascular disease. The sequences
XX are also used in drug screening assays.
XX
XX Sequence 24 BP; 7 A; 6 C; 7 G; 4 T; 0 other;
SQ Query Match 67.8%; Score 12.2; DB 22; Length 24;
Best Local Similarity 58.8%; Pred. No. 1.6e+03;
Matches 10; Conservative 4; Mismatches 3; Indels 0; Gaps 0;

QY 2 AUGGCCUAUCGGUGCGA 18
I : I I I I I I I I I I
D b 24 ATGCCCTATCGTGCTA 8

RESULT 11
ABI84452
ID ABI84452 standard; DNA; 24 BP.
AC ABI84452;
XX
XX 15-FEB-2002 (first entry)
XX
XX Capture oligonucleotide zip ID#1028 oligo #1.
XX
XX Human; K-ras; PCR primer; probe; capture probe; mutation detection;
XX ligase detection reaction; LDR; p53; BRCA1; BRCA2; infectious disease;
XX infection; 21 hydroxylase deficiency; Turner Syndrome; Obesity;
XX cancer; oncogene; tumour suppressor; human papillomavirus; forensic;
XX environmental monitoring; food industry; feed industry; ss.
XX
XX Synthetic.
XX
XX WO200179548-A2.
XX
XX 25-OCT-2001.
XX
XX 04-APR-2001; 2001WO-US10958.
XX
XX 14-APR-2000; 2000US-197271P.
XX
XX (CORR ) CORNELL RES FOUND INC.
XX
XX Barany F, Zirvi M, Gerry NP, Favis R, Kliman R;
XX WPI; 2002-034366/04.
XX
XX Designing capture oligonucleotide probes for use on a support to which
XX complementary oligonucleotides hybridize with little mismatch -
XX
XX Example 5; Fig 25; 300pp; English.
XX
XX
```

XX The present invention describes a method (M1) for designing capture
 CC oligonucleotide probes (I) for use on a support to which complementary
 CC oligonucleotide probes (II) will hybridize with little mismatch, where
 CC (I) have melting temperatures within a narrow range. The method is useful
 CC for detecting infectious diseases caused by bacterial infectious agents
 CC e.g. Salmonella, Listeria monocytogenes and Haemophilus influenza, fungal
 CC infectious agents e.g. Cryptococcus neoformans, Candida albicans and
 CC Aspergillus fumigatus, viruses e.g. T-cell lymphocytotropic virus,
 CC Epstein-Barr virus and polio virus, and parasitic infectious agents
 CC selected from Onchocerca volvulus, Entamoeba histolytica and Dracunculus
 CC medinensis. The method is also useful for detecting genetic diseases such
 CC as 21 hydroxylase deficiency, Turner Syndrome and obesity defects.
 CC Detecting cancer involving oncogenes, tumour suppressor genes, or genes
 CC involved in DNA amplification, replication, recombination or repair, the
 CC cancer is specifically associated with a gene selected from BRCA1 gene,
 CC p53 gene, human papillomavirus types 16 and 18 and liver cancers. The
 CC method is also used for environmental monitoring, forensics and the food
 CC and feed industry, detecting comprises scanning (using e.g. a scanning
 CC electron microscope and infrared microscope) the support at the
 CC particular sites and identifying if ligation of the oligonucleotide probe
 CC sets occurred and correlating (using a computer) identified ligation to a
 CC presence or absence of the target nucleotide sequences. ABI82074 to
 CC ABI97546 represent oligonucleotide sequences used in the exemplification
 CC of the present invention.

XX Sequence 24 BP; 7 A; 7 C; 5 G; 5 T; 0 other;

Query Match 67.8%; Score 12.2; DB 24; Length 24;
 Best Local Similarity 64.7%; Pred. No. 1.6e+03;
 Matches 11; Conservative 3; Mismatches 3; Indels 0; Gaps 0;

QY 2 AUGCCUAUCGGUGCGA 18
 I: | | | | |
 Db 7 ATGACCAATCGATCGCA 23

RESULT 12
 ABI84453/c
 ID ABI84453 standard; DNA; 24 BP.

XX ABI84453;

XX 15-FEB-2002 (first entry)

XX Capture oligonucleotide zip ID#1028 oligo #2.

XX Human; K-ras; PCR primer; probe: capture probe; mutation detection;
 KW ligase detection reaction; LDR; p53; BRCA1; BRCA2; infectious disease;
 KW infection; 21 hydroxylase deficiency; Turner Syndrome; obesity;
 KW cancer; oncogene; tumour suppressor; human papillomavirus; forensic;
 KW environmental monitoring; food industry; feed industry; ss.

XX Synthetic.

XX WO200179548-A2.

XX 25-OCT-2001.

XX 04-APR-2001; 2001WO-US10958.

XX 14-APR-2000; 2000US-197271P.

XX (CORR) CORNELL RES FOUND INC.

XX Barany F, Zirvi M, Gerry NP, Favis R, Kliman R;

XX WPI; 2002-034366/04.

XX Designing capture oligonucleotide probes for use on a support to which
 PT complementary oligonucleotides hybridize with little mismatch -

XX Example 5; Fig 25; 300pp; English.

XX The present invention describes a method (M1) for designing capture
 CC oligonucleotide probes (I) for use on a support to which complementary
 CC oligonucleotide probes (II) will hybridize with little mismatch, where
 CC (I) have melting temperatures within a narrow range. The method is useful
 CC for detecting infectious diseases caused by bacterial infectious agents
 CC e.g. Salmonella, Listeria monocytogenes and Haemophilus influenza, fungal
 CC infectious agents e.g. Cryptococcus neoformans, Candida albicans and
 CC Aspergillus fumigatus, viruses e.g. T-cell lymphocytotropic virus,
 CC Epstein-Barr virus and polio virus, and parasitic infectious agents
 CC selected from Onchocerca volvulus, Entamoeba histolytica and Dracunculus
 CC medinensis. The method is also useful for detecting genetic diseases such
 CC as 21 hydroxylase deficiency, Turner Syndrome and obesity defects.
 CC Detecting cancer involving oncogenes, tumour suppressor genes, or genes
 CC involved in DNA amplification, replication, recombination or repair, the
 CC cancer is specifically associated with a gene selected from BRCA1 gene,
 CC p53 gene, human papillomavirus types 16 and 18 and liver cancers. The
 CC method is also used for environmental monitoring, forensics and the food
 CC and feed industry, detecting comprises scanning (using e.g. a scanning
 CC electron microscope and infrared microscope) the support at the
 CC particular sites and identifying if ligation of the oligonucleotide probe
 CC sets occurred and correlating (using a computer) identified ligation to a
 CC presence or absence of the target nucleotide sequences. ABI82074 to
 CC ABI97546 represent oligonucleotide sequences used in the exemplification
 CC of the present invention.

XX Sequence 24 BP; 5 A; 5 C; 7 G; 7 T; 0 other;

Query Match 67.8%; Score 12.2; DB 24; Length 24;
 Best Local Similarity 64.7%; Pred. No. 1.6e+03;
 Matches 11; Conservative 3; Mismatches 3; Indels 0; Gaps 0;

QY 2 AUGCCUAUCGGUGCGA 18
 I: | | | | |
 Db 18 ATGACCAATCGATCGCA 2

RESULT 13
 AAX98255/c

ID AAX98255 standard; DNA; 29 BP.

XX AAX98255;

XX 25-OCT-1999 (first entry)

XX PCR primer used to amplify a 1659 bp fragment containing ORF1.

XX Human pathogen; virulence polypeptide; virulence factor;
 KW pathogenic infection; Pseudomonas aeruginosa infection; PCR primer; ss.

XX Synthetic.

XX Pseudomonas aeruginosa.

XX WO927129-A1.

XX 03-JUN-1999.

XX 25-NOV-1998; 98WO-US25247.

XX 25-NOV-1997; 97US-0066517.

XX (GEO) GEN HOSPITAL CORP.

XX Ausubel F, Cao H, Drenkard E, Goodman HM, Mahajan-Miklos S;

XX Rahme LG, Tan M, Tsongalis J;

XX WPI; 1999-357851/30.

XX Virulence factors useful in developing disease treatments

XX Disclosure; Page 26; 228pp; English.

XX The present sequence represents a PCR primer used to amplify Pseudomonas

CC aeruginosa nucleic acid sequences, in the course of the invention.
 CC P. aeruginosa is an opportunistic human pathogen present in soil
 CC water and plants. The specification describes virulence polypeptides
 CC and nucleic acid sequence encoding such polypeptides. These sequences
 CC can be used to identify a compound which is capable of decreasing the
 CC expression of a pathogenic virulence factor. Compounds that inhibit
 CC the expression or activity of virulence factor polypeptides can be
 CC used to treat pathogenic infections, especially where the infection
 CC is a P. aeruginosa infection.
 XX Sequence 29 BP; 6 A; 10 C; 8 G; 5 T; 0 other;
 SQ Query Match 67.8%; Score 12.2; DB 20; Length 29;
 Best Local Similarity 64.7%; Pred. No. 1.7e+03;
 Matches 11; Conservative 3; Mismatches 3; Indels 0; Gaps 0;
 QY 1 A AUGGCCUAUGGUGCG 17
 |||||:|:|:|:|:|
 DB 19 AACGGCGTATCGTGGC 3
 RESULT 14
 AAT70787/c
 ID AAT70787 standard; RNA; 40 BP.
 XX
 AC AAT70787;
 XX
 DT 30-JUL-1997 (first entry)
 XX
 DE Stenotic carotid artery binding ligand C1v19.
 XX
 KW Ligand: peripheral blood mononuclear cell; fibrin clot; carotid artery;
 KW systematic evolution of ligands by exponential enrichment method; PBMC;
 KW epitope; macromolecule; tissue SELEX method; cancer screening; therapy;
 KW AIDS monitoring; localisation of thrombi; atherosclerosis; ss.
 XX
 OS Synthetic.
 XX
 FH Key Location/Qualifiers
 FT modified_base 1..40
 FT /*tag= a
 FT /mod_base= all C's are 2-fluorine-C
 FT modified_base 1..40
 FT /*tag= b
 FT /mod_base= all U's are 2-fluorine-U
 FT
 PN W09634874-A1.
 XX
 PD 07-NOV-1996.
 XX
 PF 01-MAY-1996; 96WO-US06059.
 XX
 PR 03-MAY-1995; 95US-0433126.
 PR 03-MAY-1995; 95US-0433124.
 XX
 PA (NEXS-) NEXSTAR PHARM INC.
 PA (SCHD) SCHERING AG.
 XX
 PI Gold L, Schneider DJ, Speck U, Stephens A;
 XX
 DR WPI; 1996-506091/50.
 XX
 XX Nucleic acid ligands used in cancer screening, AIDS monitoring etc.
 PT - bind to peripheral blood mononuclear cells, fibrin clots or
 PT carotid arteries
 XX
 PS Claim 23; Page 58; 138pp; English.
 XX
 CC AAT70705-T70803 represent the random regions from a degenerate ssDNA
 CC library based on the sequence represented by AAT42819. These sequences
 CC are all ligands specific for fibrin clots identified using the method of
 CC the invention. The method of the invention is for identifying nucleic
 CC acid (NA) ligands and NA ligand sequences to a tissue target selected

CC from peripheral blood mononuclear cells (PBMC's) (such as the ligands
 CC represented by AAT70584-T70616), fibrin clots (such as the ligands
 CC represented by AAT70617-T70704), and carotid arteries (such as these
 CC ligands). The method comprises preparing a candidate mixture of NA
 CC sequences, and contacting these with the tissue, whereby NAs having an
 CC increased affinity to the tissue relative to the candidate mixture may
 CC be partitioned from the remainder of the candidate mixture. The
 CC increased affinity NAs are then partitioned from the remainder of the
 CC candidate mixture, and are amplified to yield a mixture of NAs enriched
 CC for NA sequences with a relatively higher affinity and specificity for
 CC binding to the tissue, whereby NA ligands of the tissue may be
 CC identified. The method represents a tissue SELEX (systematic evolution of
 CC ligands by exponential enrichment) method. The NA ligands and epitopes
 CC and macromolecules identified using the ligands can be used in diagnostic
 CC and therapeutic applications. In particular, they can be used for e.g.
 CC cancer screening, AIDS monitoring, detection and localisation of thrombi
 CC or atherosclerosis diagnosis and therapy.
 XX
 SQ Sequence 40 BP; 9 A; 13 C; 8 G; 10 U; 0 other;
 Query Match 67.8%; Score 12.2; DB 17; Length 40;
 Best Local Similarity 64.7%; Pred. No. 1.8e+03;
 Matches 11; Conservative 3; Mismatches 3; Indels 0; Gaps 0;
 QY 2 AUGGCCUAUGGUGCGA 18
 |||||:|:|:|:|:|
 DB 21 AAGGCCCTTCGGTCCGA 5
 |||||:|:|:|:|:|

RESULT 15
 AAT21824/c
 ID AAT21824 standard; cDNA to mRNA; 53 BP.
 XX
 AC AAT21824;
 XX
 DT 01-AUG-1996 (first entry)
 XX
 DE Human gene signature HUMGS03315.
 XX
 KW Gene signature; messenger RNA; mRNA; relative abundance; frequency;
 KW human; cloning; mapping; non-biased library; diagnosis; detection;
 KW cell typing; abnormal cell function; ss.
 XX
 OS Homo sapiens.
 XX
 PN W09514772-A1.
 XX
 PD 01-JUN-1995.
 XX
 PF 11-NOV-1994; 94WO-JP01916.
 XX
 PR 12-NOV-1993; 93JP-0355504.
 XX
 PA (MATS/) MATSUBARA K.
 PA (OKUB/) OKUBO K.
 XX
 PI Matsubara K, Okubo K;
 XX
 DR WPI; 1995-206931/27.
 XX
 XX Identifying gene signatures in 3'-directed human cDNA library - e.g.
 PT for diagnosis of abnormal cell function, by preparing cDNA that
 PT reflects relative abundance of corresp. mRNA in specific human
 PT tissues
 XX
 PS Claim 1; Page 975; 2245pp; Japanese.
 XX
 CC A single-stranded DNA (or its complementary strand or the corresp.
 CC double-stranded DNA) which comprises one of the 7837 "GS" sequences
 CC given in AAT19001-T26837 and which is able to hybridise to part of
 CC human genomic DNA, cDNA or mRNA is claimed. The GS (Gene Signature)
 CC sequences were obtained from 3'-directed cDNA libraries prepared from
 CC from various human tissues; synthesis of cDNA was initiated from the

3'-end of mRNA by using poly(T) as the sole primer. Since the 3'-untranslated sequence is unique to a particular mRNA species, almost all the 3'-oriented cDNAs hybridise with specific mRNAs. Each library is constructed so as to reflect accurately the relative abundance of different mRNAs in the particular tissue from which it was derived.

Sequence 53 BP; 14 A; 11 C; 12 G; 15 T; 1 other;

Query Match 67.8%; Score 12.2; DB 16; Length 53;
Best Local Similarity 64.7%; Pred. No. 1.8e+03;
Matches 11; Conservative 3; Mismatches 3; Indels 0; Gaps 0;

QY 1 A AUGGCCUAUCGGUGCG 17
||:||||| :|| :|||
Db 34 AATGGCCCCCTCGATGCG 18

Search completed: July 6, 2003, 14:32:53
Job time : 194.091 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: July 6, 2003, 14:26:51 ; Search time 1209.27 Seconds

(without alignments)
241.069 Million cell updates/sec

Title: us-09-780-929-98

Perfect score: 18

Sequence: 1 aauggccuauccgucga 18

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 16154066 seqs, 8097743376 residues

Total number of hits satisfying chosen parameters: 146654

Minimum DB seq length: 0

Maximum DB seq length: 60

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

EST.*

1: em_estba.*
2: em_esthum.*
3: em_estin.*
4: em_estnu.*
5: em_estov.*
6: em_estpl.*
7: em_estro.*
8: em_hic.*
9: gb_est1.*
10: gb_est2.*
11: gb_hic.*
12: gb_est3.*
13: gb_est4.*
14: gb_est5.*
15: em_estfun.*
16: em_estom.*
17: gb_gss.*
18: em_gss_hum.*
19: em_gss_inv.*
20: em_gss_pln.*
21: em_gss_vrt.*
22: em_gss_fun.*
23: em_gss_mam.*
24: em_gss_mus.*
25: em_gss_other.*
26: em_gss_pro.*
27: em_gss_rod.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	13.8	76.7	49	17	TA58C07Q
c 2	12.4	68.9	50	9	AU183461
3	12.2	67.8	37	14	H13124
c 4	12.2	67.8	50	14	C00960
5	11.8	65.6	50	9	AU105634
c 6	11.8	65.6	56	10	AV847604

7	11.6	64.4	29	17	AZ800642
8	11.6	64.4	55	9	AA863171
c 9	11.2	62.2	30	17	AZ840293
c 10	11.2	62.2	36	17	AZ605771
11	11.2	62.2	37	9	AA931624
c 12	11.2	62.2	45	17	AZ317769
13	11.2	62.2	51	17	BH618002
14	11.2	62.2	51	17	BH618002
15	11.2	62.2	52	9	AA721034
c 16	11	61.1	32	17	TA371C10P
c 17	11	61.1	38	9	A1221514
c 18	10.8	60.0	32	13	BJ066180
19	10.8	60.0	44	17	BH252563
c 20	10.8	60.0	58	9	A1953654
21	10.8	60.0	59	17	AL758565
c 22	10.6	58.9	27	17	TA208E12P
23	10.6	58.9	38	17	TA244G07P
c 24	10.6	58.9	40	9	A1613042
c 25	10.6	58.9	40	17	AZ511081
c 26	10.6	58.9	44	17	AZ495581
c 27	10.6	58.9	46	9	AA455514
c 28	10.6	58.9	50	9	AU102871
c 29	10.6	58.9	50	17	AZ777046
c 30	10.6	58.9	55	17	BH810903
31	10.6	58.9	56	14	H42612
32	10.6	58.9	58	17	BH256481
33	10.4	57.8	27	17	AQ025667
c 34	10.4	57.8	52	9	AA485733
c 35	10.2	56.7	33	12	BF026570
c 36	10.2	56.7	34	9	A1192963
37	10.2	56.7	35	17	TA177A10Q
38	10.2	56.7	36	17	BH751662
39	10.2	56.7	36	17	BH751810
40	10.2	56.7	38	17	BH611852
41	10.2	56.7	38	17	BH618406
42	10.2	56.7	38	17	BH618437
43	10.2	56.7	38	17	BH618558
44	10.2	56.7	38	17	BH751595
45	10.2	56.7	39	13	BI838507

ALIGNMENTS

RESULT 1
TA58C07Q
LOCUS
DEFINITION
T. brucei sheared genomic DNA clone 58c07, reverse sequence,
genomic survey sequence.
ACCESSION
AL455710
VERSION
AL455710.1 GI:11857988
KEYWORDS
GSS.
SOURCE
Trypanosoma brucei.
ORGANISM
Trypanosoma brucei
Eukaryota; Euglenozoa; Kinetoplastida; Trypanosomatidae;
Trypanosoma.
REFERENCE
1 (bases 1 to 49)
Hall, N., Bowman, S., Lennard, N.J., Doggett, J., Atkin, R.,
Chillingworth, C., Ormond, D., Harris, B., El-Sayed, N., Hou, L.,
Melville, S.E., Rajandream, M.A. and Barrell, B.G.
Direct Submission
Submitted (10-DEC-2000) Trypanosoma brucei genome sequencing
project, Sanger Centre, The Wellcome Trust Genome Campus, Hinxton,
Cambridge CB10 1SA, E-mail: barrellesanger.ac.uk and
nhlesanger.ac.uk
COMMENT
Constructed at the Institute for Genomic Research (TIGR),
Rockville, MD. Genomic DNA isolated from a cloned population of
Trypanosoma brucei (TREV027/4 GUFat 10.1) was mechanically sheared
to give a tight size distribution (4 kb). The v + i method used for the library construction is
described in detail in Smith, H. and Venter, J.C. (Making small
insert libraries for whole genome shotgun sequencing projects. In
Genome Sequencing: A Practical Approach, eds. M. Vaudin and B.

Barrell, Oxford University Press, 1999).

Email: nelsayed@tigr.org
Details of T. brucei sequencing at the Sanger Centre are available
at http://www.sanger.ac.uk/Projects/T_brucei/.

FEATURES

source

1..49
/organism="Trypanosoma brucei"
/strain="TREU927"
/db_xref="taxon:5691"
/clone="58c07"

BASE COUNT
ORIGIN

15 a 11 c 14 g 9 t

Query Match 76.7%; Score 13.8; DB 17; Length 49;

Best Local Similarity 70.6%; Pred. No. 7.7e+03;
Matches 12; Conservative 3; Mismatches 2; Indels 0; Gaps 0;

QY 2 AUGGCCUACUGGUGCGA 18

1: 111111111111

Db 29 ATAGCCGATCGGTGCGA 45

RESULT 2

AU183461/c

LOCUS

DEFINITION AU183461 Cyprinus carpio head kidney stimulated by
lipo-polysaccharide and concanavalin-A Cyprinus carpio cDNA clone
H119, mRNA sequence.

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

Cyprinus carpio
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Actinopterygii; Neopterygii; Teleostei; Ostariophysi; Cypriniformes

REFERENCE

AUTHORS

TITLE

Analysis of expressed sequence tags (EST) obtained from common carp
, Cyprinus carpio L., head kidney cells after stimulation by two
mitogens, lipo-polysaccharide and concanavalin-A

JOURNAL

MEDLINE

COMMENT

2160774
Contact: Masahiro Sakai
Faculty of Agriculture
Miyazaki University
1-1 nishiki gakuenkibadai, Miyazaki, Miyazaki 889-2192, Japan
Email: m.sakai@cc.miyazaki-u.ac.jp.

FEATURES

source

1..50
/organism="Cyprinus carpio"
/db_xref="taxon:7962"
/clone="H119"

/clone_lib="Cyprinus carpio head kidney stimulated by
lipo-polysaccharide and concanavalin-A"
/tissue_type="head kidney"
/note="common name: common carp; stimulated by
lipo-polysaccharide and concanavalin-A"

BASE COUNT

ORIGIN

9 a 11 c 19 g 11 t

Query Match 68.9%; Score 12.4; DB 9; Length 50;

Best Local Similarity 78.6%; Pred. No. 3.7e+04;
Matches 11; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 5 GCCUACUGGUGCGA 18

111111111111

Db 50 GCCATCGGTGCGA 37

RESULT 3

H13124

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

H13124

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

H13124

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

H13124

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

H13124

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

H13124

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

H13124

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

H13124

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

H13124

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

H13124

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

H13124

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

H13124

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

H13124

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

H13124

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

H13124

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

H13124

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

H13124

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

H13124

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

H13124

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

H13124

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

H13124

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

H13124

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
 REFERENCE 1 (bases 1 to 50)
 AUTHORS Suzuki, Y., Yoshitomo-Nakagawa, K., Maruyama, K., Suyama, A. and Sugano S.
 TITLE Construction and characterization of a full length-enriched and a 5'-end-enriched cDNA library. Gene 200 (1-2), 149-156 (1997).
 JOURNAL
 COMMENT BodyMap: human gene expression database
 Contact: Okubo, K.
 Institute for Molecular and Cellular Biol
 Osaka University
 1-3, Yamada-oka, Suita, Osaka Pref. 565, Japan
 Tel: 06-877-5111(ex.3315)
 Email: kousaku@imcb.osaka-u.ac.jp
 Human Gene Signature, 3'-directed cDNA sequence. We are not submitting the same cDNA sequence redundantly to DBJ since 1993.
 For the abundance information of clones with this sequence in this library and as well as in other 3'-directed libraries, see, <http://www.imcb.osaka-u.ac.jp/bodymap/>. The sequences of the clones represented by this GS sequences is also found there.

FEATURES source

1..50
 Location/Qualifiers
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /clone_lib="Human adult (K.Okubo)"
 /dev_stage="adult"
 /note="Organ: blood; Vector: l-gt-11; Site: Eco-RI; Monocytes were prepared from blood by ficoll-hypaque, percoll and T cell rosetting purification steps (purity: 96 %). mRNA was prepared from activated monocytes from a patient with rheumatoid arthritis. mRNA was reverse transcribed with MuLV. Using Eco-RI linkers cDNA was cloned into l-gt-11 vector arms. The cDNA library was screened by differential hybridization using radioactively marked ss-cDNA from activated and non-activated monocytes."

BASE COUNT 11 a 11 c 12 g 15 t 1 others

Query Match 67.8%; Score 12.2; DB 14; Length 50;
 Best Local Similarity 64.7%; Pred. No. 4.6e+04;
 Matches 11; Conservative 3; Mismatches 3; Indels 0; Gaps 0;

QY 1 AUGGCCUACGUGGCG 17
 ||:||||: ||: |||
 DB 34 AATGGCCCTCGATGCG 18

RESULT 5 AUI05634

LOCUS AUI05634 Sugano Homo sapiens cDNA library Homo sapiens cDNA clone
 DEFINITION KAT08740, mRNA sequence.
 ACCESSION AUI05634
 VERSION AUI05634.1 GI:13555155
 KEYWORDS EST.
 SOURCE human.

ORGANISM

Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
 REFERENCE 1 (bases 1 to 50)

AUTHORS Suzuki, Y., Taira, H., Tsunoda, T., Mizushima-Sugano, J., Sese, J., Hata S., Ota, T., Isogai, T., Tanaka, T., Morishita, S., Okubo, K., Sakaki Y., Nakamura, Y., Suyama, A. and Sugano, S.
 TITLE Diverse transcriptional initiation revealed by fine, large-scale mapping of mRNA start sites

JOURNAL EMBO Rep. 2 (5), 388-393 (2001)
 MEDLINE 21270072

COMMENT Contact: Yutaka Suzuki
 Department of Virology
 Institute of Medical Science, University of Tokyo
 4-6-1, Shirokanedai, Minatoku, Tokyo 108-8639, Japan
 Email: ysuzuki@ims.u-tokyo.ac.jp

Suzuki, Y., Yoshitomo-Nakagawa, K., Maruyama, K., Suyama, A. and Sugano S.
 Construction and characterization of a full length-enriched and a 5'-end-enriched cDNA library. Gene 200 (1-2), 149-156 (1997).

FEATURES source

1..50
 Location/Qualifiers
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /clone_lib="KAT08740"
 /clone_lib="Sugano Homo sapiens cDNA library"
 /note="Differential display comparison of untreated and dimethylfumarate treated U937 cells"

BASE COUNT 10 a 14 c 14 g 12 t

ORIGIN

Query Match 65.6%; Score 11.8; DB 9; Length 50;
 Best Local Similarity 66.7%; Pred. No. 7.2e+04;
 Matches 10; Conservative 3; Mismatches 2; Indels 0; Gaps 0;

QY 4 GGCCUACGUGGCGA 18
 |||:|:|:|:|:|

DB 20 GCGTATCCGTGCGA 34

RESULT 6 AV847604/c

LOCUS AV847604 Nori Satoh unpublished cDNA library, larva Ciona
 DEFINITION intestinalis cDNA clone rcilv08b14 3', mRNA sequence.
 ACCESSION AV847604
 VERSION AV847604.1 GI:16828139
 KEYWORDS EST.
 SOURCE Ciona intestinalis.

ORGANISM

Ciona intestinalis
 Eukaryota; Metazoa; Chordata; Urochordata; Ascidiacea; Enterogona; Phlebobranchia; Clonidae; Ciona.
 REFERENCE 1 (bases 1 to 56)
 AUTHORS Satoh, N., Satou, Y., Kohara, Y. and Shin-i, T.
 TITLE Expressed genes in Ciona intestinalis
 JOURNAL Unpublished (2000)
 COMMENT Contact: Nori Satoh
 Department of Zoology
 Kyoto University
 Sakyo-ku, Kyoto, Kyoto 606-8502, Japan
 Tel: 81-75-753-4081
 Fax: 81-75-705-1113
 Email: satoheascidian.zool.kyoto-u.ac.jp.

FEATURES source

1..56
 Location/Qualifiers
 /organism="Ciona intestinalis"
 /db_xref="taxon:7719"
 /clone_lib="rcilv08b14"
 /clone_lib="Nori Satoh unpublished cDNA library, larva"
 /tissue_type="whole animal"
 /dev_stage="larva"
 /note="vector: pBluescript SK"

BASE COUNT 12 a 11 c 18 g 15 t

ORIGIN

Query Match 65.6%; Score 11.8; DB 10; Length 56;
 Best Local Similarity 73.3%; Pred. No. 7.4e+04;
 Matches 11; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 2 AUGCCUACGUGGCG 16
 |:||||:|:|:|:|

DB 45 ATGCCCAACGGTGC 31

RESULT 7

AZ800642
 LOCUS AZ800642
 DEFINITION 2M005861R Mouse 10kb plasmid UUGClm library Mus musculus genomic clone UUGC2M005861R, DNA sequence.
 ACCESSION AZ800642

```

VERSION      AZ800642.1  GI:12952965
KEYWORDS     GSS.
SOURCE       Mus musculus
ORGANISM     house mouse.

REFERENCE    Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
AUTHORS      Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
              1 (bases 1 to 29)
              Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C.,
              Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly
              M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A.
              and Wright,D., Weiss,R.

TITLE        Mouse whole genome scaffolding with paired end reads from 10kb
              plasmid inserts

JOURNAL      Unpublished (2000)
COMMENT      Contact: Robert B. Weiss
              University of Utah Genome Center
              Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT
              84112, USA
              Tel: 801 585 5606
              Fax: 801 585 7177
              Email: ddunn@genetics.utah.edu
              Insert Length: 10000 Std Error: 0.00
              Plate: 0058 row: G column: 16
              Seq primer: CACACAGGAAACAGCTATGACC
              Class: plasmid ends
              High quality sequence stop: 29.

FEATURES     Location/Qualifiers
              source          1..29
              /organism="Mus musculus"
              /db_xref="taxon:10090"
              /clone="UUGC2M0058G16"
              /clone_lib="Mouse 10kb plasmid UUGCLM library"
              /sex="Male"
              /lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
              /note="Vector: PWD42nv; Purified genomic DNA from M.
              musculus C57BL/6J (male) was obtained from the Jackson
              Laboratory Mouse DNA Resource
              (http://www.jax.org/resources/documents/dnares/). The DNA
              was hydrodynamically sheared by repeated passage through a
              0.005 inch orifice at constant velocity. The sheared DNA
              was blunt end-repaired with T4 DNA polymerase and T4
              polynucleotide kinase. Adaptor oligonucleotides were
              ligated to the blunt ends in high molar excess. The
              adaptor DNA was purified and size-selected for a 9.5 to
              10.5 kb range using preparative agarose gel
              electrophoresis. Vector DNA was prepared from a derivative
              of PWD42 (gi14732114|gb|AF129072.1), a copy-number
              inducible derivative of plasmid R1. The vector was ligated
              with adaptors complementary to the insert adaptors and
              purified. The sheared, adaptor mouse DNA was annealed to
              adaptor vector DNA, and transformed into
              chemically-competent E. coli XL10-Gold (Stratagene) cells
              and selected for ampicillin resistance."
BASE COUNT   4 a 5 c 7 g 13 t
ORIGIN
Query Match      64.4%; Score 11.6; DB 17; Length 29;
Best Local Similarity 61.1%; Pred. No. 7.8e+04;
Matches 11; Conservative 3; Mismatches 4; Indels 0; Gaps 0;

Qy 1 AAUGGCCUACUGGUGCGCA 18
    |||||:|:|:|:|
Db 2 ATTGCGCTATGGGGCGCA 19

RESULT 8
AA863171
LOCUS      55 bp mRNA linear EST 29-APR-1998
DEFINITION o91e02.s1 NCI-CCAP_Ki5 Homo sapiens cDNA clone IMAGE:1455674 3'
            similar to SW:H1E3_HUMAN Q15738 H105E3 PROTEIN. ; mRNA sequence.
ACCESSION  AA863171

```

```

VERSION      AA863171.1  GI:2955650
KEYWORDS     EST.
SOURCE       human.
ORGANISM     Homo sapiens
              Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
              Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
              1 (bases 1 to 55)
              NCI-CCAP http://www.ncbi.nlm.nih.gov/ncicgap.
              National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
              Tumor Gene Index
              Unpublished (1997)
              COMMENT      Contact: Robert Strausberg, Ph.D.
              Email: cgaps-r@mail.nih.gov
              Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R.
              Emert-Buck, M.D., Ph.D.
              CDNA Library Preparation: M. Bento Soares, Ph.D.
              CDNA Library Arrayed by: Greg Lennon, Ph.D.
              DNA Sequencing by: Washington University Genome Sequencing Center
              Clone distribution: NCI-CCAP clone distribution information can be
              found through the I.M.A.G.E. Consortium/LLNL at:
              www-bio.llnl.gov/bbrp/image/image.html

Trace considered overall poor quality
Insert Length: 360 Std Error: 0.00
Seq primer: -40ml3 fwd. ET from Amersham
High quality sequence stop: 1.

FEATURES     Location/Qualifiers
              source          1..55
              /organism="Homo sapiens"
              /db_xref="taxon:9606"
              /clone="IMAGE:1455674"
              /clone_lib="NCI-CCAP_Ki5"
              /tissue_type="2 pooled tumors (clear cell type)"
              /lab_host="DH10B"
              /note="Organ: kidney; Vector: pT7T3D-Pac (Pharmacia) with
              a modified polylinker; Site_1: Not I; Site_2: Eco RI; 1st
              strand cDNA was primed with a Not I - oligo(dT) primer [5'
              AACTGGAGAAATTCGGCGCGCAATATATTTTATTTT 3'],
              double-stranded cDNA was ligated to Eco RI adaptors
              (Pharmacia), digested with Not I and cloned into the Not I
              and Eco RI sites of the modified pT7T3 vector. Library
              went through one round of normalization. Library
              constructed by Bento Soares and M. Fatima Bonaldo. "
BASE COUNT   12 a 16 c 15 g 12 t
ORIGIN
Query Match      64.4%; Score 11.6; DB 9; Length 55;
Best Local Similarity 66.7%; Pred. No. 9.2e+04;
Matches 12; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

Qy 1 AAUGGCCUACUGGUGCGCA 18
    |||||:|:|:|:|
Db 28 AATTGCCAACACGATGCGA 45

RESULT 9
AZ840293/c
LOCUS      30 bp DNA linear GSS 20-FEB-2001
DEFINITION 2M0136H17R Mouse 10kb plasmid UUGCLM library Mus musculus genomic
              clone UUGC2M0136H17 R, DNA sequence.
ACCESSION  AZ840293
VERSION    AZ840293.1  GI:13010201
KEYWORDS   GSS.
SOURCE     house mouse.
ORGANISM   Mus musculus
              Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
              Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
              1 (bases 1 to 30)
              Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C.,
              Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly
              M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A.
              and Wright,D., Weiss,R.

REFERENCE    Mouse whole genome scaffolding with paired end reads from 10kb
              plasmid inserts
AUTHORS

```

JOURNAL
COMMENT

plasmid inserts
Unpublished (2000)
Contact: Robert B. Weiss
University of Utah Genome Center
University of Utah
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT
84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0136 row: H column: 17
Seq primer: CACACGGAACACCTATGACC
Class: plasmid ends
High quality sequence stop: 30.

FEATURES
source

1. .30
Location/Qualifiers
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUC2M0136H17"
/clone_lib="Mouse 10kb plasmid UUC1M library"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
/note="Vector: PWD42nv; Purified genomic DNA from M.
musculus C57BL/6J (male) was obtained from the Jackson
Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA
was hydrodynamically sheared by repeated passage through a
0.005 inch orifice at constant velocity. The sheared DNA
was blunt end-repaired with T4 DNA polymerase and T4
polynucleotide kinase. Adaptor oligonucleotides were
ligated to the blunt ends in high molar excess. The
adapted DNA was purified and size-selected for a 9.5 to
10.5 kb range using preparative agarose gel
electrophoresis. Vector DNA was prepared from a derivative
of pWD42 (gi14732114|gb|AF129072.1), a copy-number
inducible derivative of plasmid R1. The vector was ligated
with adaptors complementary to the insert adaptors and
purified. The sheared, adapted mouse DNA was annealed to
adapted vector DNA, and transformed into
chemically-competent E. coli XL10-Gold (Stratagene) cells
and selected for ampicillin resistance."

BASE COUNT
ORIGIN

4 a 10 c 9 g 7 t

Query Match 62.2%; Score 11.2; DB 17; Length 30;
Best Local Similarity 62.5%; Pred. No. 1.2e+05;
Matches 10; Conservative 3; Mismatches 3; Indels 0; Gaps 0;

Qy 1 AAUGGCCUUAUGCGUGC 16
||:||||: ||||
Db 30 AATGGCTGTCCGAGC 15

RESULT 10
AZ605771/c

LOCUS
DEFINITION
clone UUC1M0427K13 F, DNA sequence.
ACCESSION
AZ605771
VERSION
AZ605771.1 GI:11727961
KEYWORDS
GSS.
SOURCE
Mus musculus
house mouse.

ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 36)
REFERENCE
AUTHORS
Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamill, C.,
Islam, H., Longacre, S., Mahmood, M., Meenen, E., Pedersen, T., Reilly,
M., Rose, M., Rose, R., Stokes, R., Tingey, A., von Niederhausern, A.
and Wright, D., Weiss, R.
TITLE
Mouse whole genome scaffolding with paired end reads from 10kb

JOURNAL
COMMENT

plasmid inserts
Unpublished (2000)
Contact: Robert B. Weiss
University of Utah Genome Center
University of Utah
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT
84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0427 row: K column: 13
Seq primer: CGTTGTAACGACGCCAGT
Class: plasmid ends
High quality sequence stop: 36.

FEATURES
source

1. .36
Location/Qualifiers
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUC1M0427K13"
/clone_lib="Mouse 10kb plasmid UUC1M library"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
/note="Vector: PWD42nv; Purified genomic DNA from M.
musculus C57BL/6J (male) was obtained from the Jackson
Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA
was hydrodynamically sheared by repeated passage through a
0.005 inch orifice at constant velocity. The sheared DNA
was blunt end-repaired with T4 DNA polymerase and T4
polynucleotide kinase. Adaptor oligonucleotides were
ligated to the blunt ends in high molar excess. The
adapted DNA was purified and size-selected for a 9.5 to
10.5 kb range using preparative agarose gel
electrophoresis. Vector DNA was prepared from a derivative
of pWD42 (gi14732114|gb|AF129072.1), a copy-number
inducible derivative of plasmid R1. The vector was ligated
with adaptors complementary to the insert adaptors and
purified. The sheared, adapted mouse DNA was annealed to
adapted vector DNA, and transformed into
chemically-competent E. coli XL10-Gold (Stratagene) cells
and selected for ampicillin resistance."

BASE COUNT
ORIGIN

14 a 10 c 5 g 7 t

Query Match 62.2%; Score 11.2; DB 17; Length 36;
Best Local Similarity 56.2%; Pred. No. 1.3e+05;
Matches 9; Conservative 4; Mismatches 3; Indels 0; Gaps 0;

Qy 2 AUGGCCUUAUGCGUGCG 17
||:||||: ||||
Db 18 ATGGCTATCAGTGTG 3

RESULT 11
AA931624

LOCUS
DEFINITION
clone UUC1M0427K13 F, DNA sequence.
ACCESSION
AA931624
VERSION
AA931624.1 GI:3086010
KEYWORDS
EST.
SOURCE
human.
ORGANISM
Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
1 (bases 1 to 37)
REFERENCE
AUTHORS
NCI-CCGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
JOURNAL
Unpublished (1997)
CONTACT: Robert Strausberg, Ph.D.

Email: cgapbs-re@mail.nih.gov
 Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R. Emmert-Buck, M.D., Ph.D.
 cDNA Library Preparation: M. Bento Soares, Ph.D.
 DNA Sequencing by: Washington University Genome Sequencing Center
 Clone distribution: NCI-CGAP clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: www-bio.llnl.gov/bbrp/image/image.html

Trace considered overall poor quality
 Seq primer: ~40ml3 fwd. ET from Amersham
 High quality sequence stop: 1.

FEATURES

source

1. 37
 Location/Qualifiers
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /clone="IMAGE:1568151"
 /clone_lib="NCI-CGAP_Lu5"
 /tissue_type="carcinoid"
 /lab_host="DH10B"
 /note="Organ: lung; Vector: pT7T3D-Pac (Pharmacia) with a modified polylinker; 1st strand cDNA was prepared from neuroendocrine lung carcinoid, and was then primed with a Not I - oligo(dT) primer. Double-stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of the modified pT7T3 vector. Library is normalized. Library was constructed by Bento Soares and M. Fatima Bonaldo."

BASE COUNT

ORIGIN

7 a 10 c 11 g 9 t

Query Match 62.2%; Score 11.2; DB 9; Length 37;

Best Local Similarity 56.2%; Pred. No. 1.3e+05;

Matches 9; Conservative 4; Mismatches 3; Indels 0; Gaps 0;

QY 1 AAUGGCCUUAUGGUGC 16

DB 20 AATTGCCTATGGGTCC 35

RESULT 12

AZ317769/c

LOCUS

DEFINITION 45 bp DNA linear GSS 29-SEP-2000

clone UUGCLM0036002 R, DNA sequence.

ACCESSION AZ317769

VERSION AZ317769.1 GI:10366900

KEYWORDS GSS.

SOURCE house mouse.

ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus.

Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C.,

Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly

,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A.

and Wright,D., Weiss,R.

Mouse whole genome scaffolding with paired end reads from 10kb

plasmid inserts

Unpublished (2000)

Contact: Robert B. Weiss

University of Utah Genome Center

University of Utah

Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLG, UT

84112, USA

Tel: 801 585 5606

Fax: 801 585 7177

Email: ddunn@genetics.utah.edu

Insert Length: 10000 Std Error: 0.00

Plate: 0036 row: 0 column: 02

Seq primer: CACACAGGAACAGCTATGACC

Class: plasmid ends

High quality sequence stop: 45.

FEATURES

source

Location/Qualifiers

1. 45

/organism="Mus musculus"

/strain="C57BL/6J"

/db_xref="taxon:10090"

/clone="UUGCLM0036002"

/clone_lib="Mouse 10kb plasmid UUGCLM library"

/sex="Male"

/lab_host="E. Coli strain XL10-Gold, Tl-resistant, F-"

/note="Vector: PWD42nv; Purified genomic DNA from M.

musculus C57BL/6J (male) was obtained from the Jackson

Laboratory Mouse DNA Resource

(http://www.jax.org/resources/documents/dnares/). The DNA

was hydrodynamically sheared by repeated passage through a

0.005 inch orifice at constant velocity. The sheared DNA

was blunt end-repaired with T4 DNA polymerase and T4

polynucleotide kinase. Adaptor oligonucleotides were

ligated to the blunt ends in high molar excess. The

adapted DNA was purified and size-selected for a 9.5 to

10.5 kb range using preparative agarose gel

electrophoresis. Vector DNA was prepared from a derivative

of pWD42 (gil4732114|gb|AF129072.1), a copy-number

inducible derivative of plasmid R1. The vector was ligated

with adaptors complementary to the insert adaptors and

purified. The sheared, adapted mouse DNA was annealed to

adapted vector DNA, and transformed into

chemically-competent E. coli XL10-Gold (Stratagene) cells

and selected for ampicillin resistance."

BASE COUNT 10 a 11 c 12 g 12 t

ORIGIN

Query Match 62.2%; Score 11.2; DB 17; Length 45;

Best Local Similarity 56.2%; Pred. No. 1.4e+05;

Matches 9; Conservative 4; Mismatches 3; Indels 0; Gaps 0;

QY 1 AAUGGCCUUAUGGUGC 16

DB 37 AATGCTCTATCGATGC 22

RESULT 13

BH618002

LOCUS

DEFINITION 51 bp DNA linear GSS 30-JAN-2002

SALK_038348 Arabidopsis thaliana TDNA insertion lines Arabidopsis

thaliana genomic clone SALK_038348, DNA sequence.

ACCESSION BH618002

VERSION BH618002.1 GI:18428097

KEYWORDS GSS.

SOURCE thale cress.

ORGANISM

Arabidopsis thaliana

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;

Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;

Rosidae; eurosids II; Brassicales; Brassicaceae; Arabidopsi.

1 (bases 1 to 51)

Alonso,J.M., Leisse,T.J., Barajas,P., Chen,H., Cheuk,R., Gadrinab

,C., Jeske,A., Karnes,M., Kim,C.J., Parker,H., Prednis,L., Shinn,P.

, Zimmerman,J. and Ecker,J.R.

A Sequence-Indexed Library of Insertion Mutations in the

Arabidopsis Genome

Unpublished (2001)

Contact: Joseph R. Ecker

Salk Institute Genomic Analysis Laboratory (SIGNAL)

The Salk Institute for Biological Studies

10010 N. Torrey Pines Road, La Jolla, CA 92037, USA

Tel: 858 453 4100 x1752

Fax: 858 558 6379

Email: ecker@salk.edu

This is single pass sequence recovered from the left border of

TDNA.

Class: TDNA tagged.

Location/Qualifiers

1. .51

FEATURES

source

```

/organism="Arabidopsis thaliana"
/strain="Columbia 0"
/db_xref="taxon:3702"
/clone="SALK_038348"
/clone_lib="Arabidopsis thaliana TDNA insertion lines"
/note="PCR was performed on Arabidopsis thaliana lines
each of which contains one or more TDNA insertion
elements. The resultant fragment for each line was
directly sequenced to determine the genomic sequence at
the site of insertion. Details of the protocols used can
be found at http://signal.salk.edu/tdna_protocols.html"

BASE COUNT      10 a      15 c      17 g      9 t
ORIGIN

Query Match      62.2%; Score 11.2; DB 17; Length 51;
Best Local Similarity 56.2%; Pred. No. 1.4e+05;
Matches 9; Conservative 4; Mismatches 3; Indels 0; Gaps 0;

Qy 3 UGCCCUAUCGUGCGA 18
Db 29 TGCCCTATAGTGGGA 44

RESULT 14
AA721034
LOCUS      52 bp      mRNA      linear      EST 22-JAN-1998
DEFINITION nx89h04.s1 NCI_CGAP_GCB1 Homo sapiens cDNA clone IMAGE:1269463 3'
similar to WP:F56DI.3 CE01971 ;, mRNA sequence.
ACCESSION  AA721034
VERSION    AA721034.1 GI:2737169
KEYWORDS   EST.
SOURCE     human.
ORGANISM   Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE  1 (bases 1 to 52)
AUTHORS   NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE     National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
JOURNAL   Unpublished (1997)
COMMENT   Contact: Robert Strausberg, Ph.D.
Email: rcgaps-r@mail.nih.gov
Tissue Procurement: Louis M. Staudt, M.D., Ph.D., David Allman,
Ph.D., Gerald Marti, M.D.
CDNA Library Preparation: M. Bento Soares, Ph.D., M. Fatima
Bonaldo, Ph.D.
CDNA Library Arrayed by: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
www-bio.llnl.gov/bbrp/image/image.html
Insert Length: 730 Std Error: 0.00
Seq primer: -40ml3 fwd. ET from Amersham
High quality sequence stop: 1.
FEATURES
source
1..52
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:1269463"
/clone_lib="NCI_CGAP_GCB1"
/tissue_type="germinal center B cell"
/lab_host="DH10B"
/note="vector: pT73b-Pac (Pharmacia) with a modified
polylinker; Site_1: Not I; Site_2: Eco RI; 1st strand cDNA
was prepared from human tonsillar cells enriched for
germinal center B cells by flow sorting (CD20+, Igu-),
provided by Dr. Louis M. Staudt (NCI), Dr. David Allman
(NCI) and Dr. Gerald Marti (CBER). cDNA synthesis was
primed with a Not I - oligo(dT) primer
[5'-TGTTACCACTCTCAAGTGGGAGCGCGCTCAATTTTCTTTT-3'
]. Double-stranded cDNA was ligated to Eco RI adaptors
(Pharmacia), digested with Not I and cloned into the Not I
and Eco RI sites of the modified pT73 vector. Library

```

```

went through one round of normalization, and was
constructed by Bento Soares and M. Fatima Bonaldo."

BASE COUNT      12 a      12 c      21 g      7 t
ORIGIN

Query Match      62.2%; Score 11.2; DB 9; Length 52;
Best Local Similarity 68.8%; Pred. No. 1.4e+05;
Matches 11; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

Qy 1 A AUGGCCUACGUGGC 16
Db 12 AACGCCGATTGGTGC 27

RESULT 15
BH641243
LOCUS      54 bp      DNA      linear      GSS 14-FEB-2002
DEFINITION 1008046D05.2EL_y1 1008 - RescueMu Grid I Zea mays genomic, DNA
sequence.
ACCESSION  BH641243
VERSION    BH641243.1 GI:18668929
KEYWORDS   GSS.
SOURCE     Zea mays.
ORGANISM   Zea mays
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACC
clade; Panicoideae; Andropogoneae; Zea.
REFERENCE  1 (bases 1 to 54)
AUTHORS   Walbot.V.
TITLE     Maize genomic sequences found using engineered RescueMu transposon
JOURNAL   Unpublished (2001)
COMMENT   Contact: Walbot V
Department of Biological Sciences
Stanford University
855 California Ave, Palo Alto, CA 94304, USA
Tel: 650 723 2227
Fax: 650 725 8221
Email: walbot@stanford.edu
Possible ligation site of ends cut by 2 different endonucleases.
Reverse complemented post-ligation sequence from source sequence.
Plate: 1008046 row: 17
Class: transposon-tagged
Location/Qualifiers
1..54
/organism="Zea mays"
/cultivar="mixed background W23/A188/B73"
/db_xref="taxon:4577"
/clone_lib="1008 - RescueMu Grid I"
/tissue_type="leaf"
/dev_stage="adult"
/lab_host="DH10B"
/note="Organ: leaf; Vector: RescueMu (engineered from
pBluescript backbone); Site_1: BamHI; Site_2: BglII;
RescueMu is a 4.9 kb, modified maize Mu transposon
designed to allow plasmid rescue from total genomic DNA.
Mu elements insert preferentially into transcription
units. For more information on RescueMu, go to the web
site www.zmdb.iastate.edu and follow the links for
'RescueMu.' Grid I was grown at Berkeley in 2001. DNA was
extracted from leaf punches, double digested using BamHI
and BglII, and ligated to form circular plasmids. DH10B
cells were transformed and then screened on LB plates with
ampicillin."

BASE COUNT      12 a      12 c      17 g      13 t
ORIGIN

Query Match      62.2%; Score 11.2; DB 17; Length 54;
Best Local Similarity 68.8%; Pred. No. 1.4e+05;
Matches 11; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

Qy 2 AUGGCCUACGUGGC 17
Db 31 ATGGCCTTACGGGCG 46

```

Search completed: July 6, 2003, 15:28:29
Job time : 1212.27 secs

```
;
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: OBLON, SPIVAK, MCCLELLAND, MAIER & NEUSTADT,
; ADDRESSEE: P.C.
; STREET: 1755 Jefferson Davis Highway, Fourth Floor
; CITY: Arlington
; STATE: Virginia
; ZIP: 22202
;
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/07/997,133
; FILING DATE: 28-DEC-1992
; CLASSIFICATION: 530
;
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US/07/642,755
; FILING DATE: 18-JAN-1991
; ATTORNEY/AGENT INFORMATION:
; NAME: Oblon, No. 528885man F.
; REGISTRATION NUMBER: 24,618
; REFERENCE/DOCKET NUMBER: 769-226-0
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (703)521-4500
; TELEFAX: (703)486-2347
; TELEX: 248855 OPAT UR
; INFORMATION FOR SEQ ID NO: 5:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 30 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: unknown
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; US-09-997-133-5

Query Match 65.6%; Score 11.8; DB 1; Length 30;
Best Local Similarity 73.3%; Pred. No. 4.4e+02;
Matches 11; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 2 AUGGCCUAUCGGUGC 16
Db 1 ACGGCCTAGCGGTGC 15

RESULT 3
US-09-997-133-5
; Sequence 5, Application US/07997133
; GENERAL INFORMATION:
; APPLICANT: Beigonzoni, Laura
; APPLICANT: Mazue, Guy
; APPLICANT: Isacchi, Antonella
; APPLICANT: Roncucci, Romeo
; APPLICANT: Sarmientos, Paolo
; TITLE OF INVENTION: Extracellular Form of the Human
; TITLE OF INVENTION: Fibroblast Growth Factor Receptor
; NUMBER OF SEQUENCES: 8
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: OBLON, SPIVAK, MCCLELLAND, MAIER & NEUSTADT,
; ADDRESSEE: P.C.
; STREET: 1755 Jefferson Davis Highway, Fourth Floor
; CITY: Arlington
; STATE: Virginia
; ZIP: 22202
;
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/07/997,133
; FILING DATE: 28-DEC-1992
; CLASSIFICATION: 530
```

```
;
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US/07/642,755
; FILING DATE: 18-JAN-1991
; ATTORNEY/AGENT INFORMATION:
; NAME: Oblon, Norman F.
; REGISTRATION NUMBER: 24,618
; REFERENCE/DOCKET NUMBER: 769-226-0
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (703)521-4500
; TELEFAX: (703)486-2347
; TELEX: 248855 OPAT UR
; INFORMATION FOR SEQ ID NO: 5:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 30 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: unknown
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; US-09-997-133-5

Query Match 65.6%; Score 11.8; DB 5; Length 30;
Best Local Similarity 73.3%; Pred. No. 4.4e+02;
Matches 11; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 2 AUGGCCUAUCGGUGC 16
Db 1 ACGGCCTAGCGGTGC 15

RESULT 4
US-09-626-929-3
; Sequence 3, Application US/09626929
; Patent No. 6319714
; GENERAL INFORMATION:
; APPLICANT: CRAMERI, ANDREAS
; APPLICANT: STEMMER, WILLEM P.C.
; APPLICANT: MINSHULL, JEREMY
; APPLICANT: BASS, STEVEN H.
; APPLICANT: WELCH, MARK
; APPLICANT: NESS, JON E.
; APPLICANT: GUSTAFSSON, CLAES
; APPLICANT: PATTEN, PHILIP A.
; TITLE OF INVENTION: OLIGONUCLEOTIDE MEDIATED NUCLEIC ACID RECOMBINATION
; FILE REFERENCE: 02-029620US
; CURRENT APPLICATION NUMBER: US/09/626,929
; 2000-07-27
; CURRENT FILING DATE: 2000-07-27
; PRIOR APPLICATION NUMBER: 09/408,392
; PRIOR FILING DATE: 1999-09-28
; PRIOR APPLICATION NUMBER: 60/118,813
; PRIOR FILING DATE: 1999-02-05
; PRIOR APPLICATION NUMBER: 60/141,049
; PRIOR FILING DATE: 1999-06-24
; NUMBER OF SEQ ID NOS: 26
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 3
; LENGTH: 40
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Bridging
; OTHER INFORMATION: oligonucleotides
; US-09-626-929-3

Query Match 65.6%; Score 11.8; DB 4; Length 40;
Best Local Similarity 60.0%; Pred. No. 4.6e+02;
Matches 9; Conservative 4; Mismatches 2; Indels 0; Gaps 0;

QY 1 AAUGGCCUAUCGGUG 15
Db 6 ACTGGCGTATCGGTG 20
```

```
RESULT 5
US-09-484-850-3
; Sequence 3, Application US/09484850
; Patent No. 6368861
; GENERAL INFORMATION:
; APPLICANT: CRAMERI, ANDREAS
; APPLICANT: STEMMER, WILLEM P.C.
; APPLICANT: MINSHULL, JEREMY
; APPLICANT: BASS, STEVEN H.
; APPLICANT: WELCH, MARK
; APPLICANT: NESS, JON E.
; APPLICANT: GUSTAFSSON, CLAES
; APPLICANT: PATTEN, PHILLIP A.
; TITLE OF INVENTION: OLIGONUCLEOTIDE MEDIATED NUCLEIC ACID RECOMBINATION
; FILE REFERENCE: 02-029620US
; CURRENT APPLICATION NUMBER: US/09/484,850
; CURRENT FILING DATE: 2000-01-18
; PRIOR APPLICATION NUMBER: 09/408,392
; PRIOR FILING DATE: 1999-09-28
; PRIOR APPLICATION NUMBER: 60/118,813
; PRIOR FILING DATE: 1999-02-05
; PRIOR APPLICATION NUMBER: 60/141,049
; PRIOR FILING DATE: 1999-06-24
; NUMBER OF SEQ ID NOS: 26
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 3
; LENGTH: 40
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Bridging
; OTHER INFORMATION: oligonucleotides
US-09-484-850-3

Query Match      65.6%; Score 11.8; DB 4; Length 40;
Best Local Similarity 60.0%; Pred. No. 4.6e+02;
Matches 9; Conservative 4; Mismatches 2; Indels 0; Gaps 0;

QY      1 AAUGGCCUAUCGGUG 15
        | :||| :|||:|
DB      6 ACTGGCGTATCGGTG 20

RESULT 6
US-09-408-392-3
; Sequence 3, Application US/09408392
; Patent No. 6376246
; GENERAL INFORMATION:
; APPLICANT: CRAMERI, ANDREAS
; APPLICANT: STEMMER, WILLEM P.C.
; APPLICANT: MINSHULL, JEREMY
; APPLICANT: BASS, STEVEN H.
; APPLICANT: WELCH, MARK
; APPLICANT: NESS, JON E.
; APPLICANT: GUSTAFSSON, CLAES
; APPLICANT: PATTEN, PHILLIP A.
; TITLE OF INVENTION: OLIGONUCLEOTIDE MEDIATED NUCLEIC ACID RECOMBINATION
; FILE REFERENCE: 02-029620US
; CURRENT APPLICATION NUMBER: US/09/408,392
; CURRENT FILING DATE: 1999-09-28
; PRIOR APPLICATION NUMBER: 60/118,813
; PRIOR FILING DATE: 1999-02-05
; PRIOR APPLICATION NUMBER: 60/141,049
; PRIOR FILING DATE: 1999-06-24
; NUMBER OF SEQ ID NOS: 26
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 3
; LENGTH: 40
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Bridging
; OTHER INFORMATION: oligonucleotides
US-09-408-392-3

Query Match      65.6%; Score 11.8; DB 4; Length 40;
Best Local Similarity 60.0%; Pred. No. 4.6e+02;
Matches 9; Conservative 4; Mismatches 2; Indels 0; Gaps 0;

QY      1 AAUGGCCUAUCGGUG 15
        | :||| :|||:|
DB      6 ACTGGCGTATCGGTG 20

RESULT 7
US-09-626-930-3
; Sequence 3, Application US/09626930
; Patent No. 6423542
; GENERAL INFORMATION:
; APPLICANT: CRAMERI, ANDREAS
; APPLICANT: STEMMER, WILLEM P.C.
; APPLICANT: MINSHULL, JEREMY
; APPLICANT: BASS, STEVEN H.
; APPLICANT: WELCH, MARK
; APPLICANT: NESS, JON E.
; APPLICANT: GUSTAFSSON, CLAES
; APPLICANT: PATTEN, PHILLIP A.
; TITLE OF INVENTION: OLIGONUCLEOTIDE MEDIATED NUCLEIC ACID RECOMBINATION
; FILE REFERENCE: 02-029620US
; CURRENT APPLICATION NUMBER: US/09/626,930
; CURRENT FILING DATE: 2000-07-27
; PRIOR APPLICATION NUMBER: 09/408,392
; PRIOR FILING DATE: 1999-09-28
; PRIOR APPLICATION NUMBER: 60/118,813
; PRIOR FILING DATE: 1999-02-05
; PRIOR APPLICATION NUMBER: 60/141,049
; PRIOR FILING DATE: 1999-06-24
; NUMBER OF SEQ ID NOS: 26
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 3
; LENGTH: 40
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Bridging
; OTHER INFORMATION: oligonucleotides
US-09-626-930-3

Query Match      65.6%; Score 11.8; DB 4; Length 40;
Best Local Similarity 60.0%; Pred. No. 4.6e+02;
Matches 9; Conservative 4; Mismatches 2; Indels 0; Gaps 0;

QY      1 AAUGGCCUAUCGGUG 15
        | :||| :|||:|
DB      6 ACTGGCGTATCGGTG 20

RESULT 8
US-09-626-528-3
; Sequence 3, Application US/09626528
; Patent No. 6426224
; GENERAL INFORMATION:
; APPLICANT: CRAMERI, ANDREAS
; APPLICANT: STEMMER, WILLEM P.C.
; APPLICANT: MINSHULL, JEREMY
; APPLICANT: BASS, STEVEN H.
; APPLICANT: WELCH, MARK
; APPLICANT: NESS, JON E.
; APPLICANT: GUSTAFSSON, CLAES
; APPLICANT: PATTEN, PHILLIP A.
; TITLE OF INVENTION: OLIGONUCLEOTIDE MEDIATED NUCLEIC ACID RECOMBINATION
; FILE REFERENCE: 02-029620US
; CURRENT APPLICATION NUMBER: US/09/626,528
; CURRENT FILING DATE: 2000-07-27
; PRIOR APPLICATION NUMBER: 09/408,392
; PRIOR FILING DATE: 1999-09-28
```

```
US-09-408-392-3

Query Match      65.6%; Score 11.8; DB 4; Length 40;
Best Local Similarity 60.0%; Pred. No. 4.6e+02;
Matches 9; Conservative 4; Mismatches 2; Indels 0; Gaps 0;

QY      1 AAUGGCCUAUCGGUG 15
        | :||| :|||:|
DB      6 ACTGGCGTATCGGTG 20

RESULT 7
US-09-626-930-3
; Sequence 3, Application US/09626930
; Patent No. 6423542
; GENERAL INFORMATION:
; APPLICANT: CRAMERI, ANDREAS
; APPLICANT: STEMMER, WILLEM P.C.
; APPLICANT: MINSHULL, JEREMY
; APPLICANT: BASS, STEVEN H.
; APPLICANT: WELCH, MARK
; APPLICANT: NESS, JON E.
; APPLICANT: GUSTAFSSON, CLAES
; APPLICANT: PATTEN, PHILLIP A.
; TITLE OF INVENTION: OLIGONUCLEOTIDE MEDIATED NUCLEIC ACID RECOMBINATION
; FILE REFERENCE: 02-029620US
; CURRENT APPLICATION NUMBER: US/09/626,930
; CURRENT FILING DATE: 2000-07-27
; PRIOR APPLICATION NUMBER: 09/408,392
; PRIOR FILING DATE: 1999-09-28
; PRIOR APPLICATION NUMBER: 60/118,813
; PRIOR FILING DATE: 1999-02-05
; PRIOR APPLICATION NUMBER: 60/141,049
; PRIOR FILING DATE: 1999-06-24
; NUMBER OF SEQ ID NOS: 26
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 3
; LENGTH: 40
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Bridging
; OTHER INFORMATION: oligonucleotides
US-09-626-930-3

Query Match      65.6%; Score 11.8; DB 4; Length 40;
Best Local Similarity 60.0%; Pred. No. 4.6e+02;
Matches 9; Conservative 4; Mismatches 2; Indels 0; Gaps 0;

QY      1 AAUGGCCUAUCGGUG 15
        | :||| :|||:|
DB      6 ACTGGCGTATCGGTG 20

RESULT 8
US-09-626-528-3
; Sequence 3, Application US/09626528
; Patent No. 6426224
; GENERAL INFORMATION:
; APPLICANT: CRAMERI, ANDREAS
; APPLICANT: STEMMER, WILLEM P.C.
; APPLICANT: MINSHULL, JEREMY
; APPLICANT: BASS, STEVEN H.
; APPLICANT: WELCH, MARK
; APPLICANT: NESS, JON E.
; APPLICANT: GUSTAFSSON, CLAES
; APPLICANT: PATTEN, PHILLIP A.
; TITLE OF INVENTION: OLIGONUCLEOTIDE MEDIATED NUCLEIC ACID RECOMBINATION
; FILE REFERENCE: 02-029620US
; CURRENT APPLICATION NUMBER: US/09/626,528
; CURRENT FILING DATE: 2000-07-27
; PRIOR APPLICATION NUMBER: 09/408,392
; PRIOR FILING DATE: 1999-09-28
```

;; PRIOR APPLICATION NUMBER: 60/118,813
;; PRIOR FILING DATE: 1999-02-05
;; PRIOR APPLICATION NUMBER: 60/141,049
;; PRIOR FILING DATE: 1999-06-24
;; NUMBER OF SEQ ID NOS: 26
;; SOFTWARE: PatentIn ver. 2.1
;; SEQ ID NO 3
;; LENGTH: 40
;; TYPE: DNA
;; ORGANISM: Artificial Sequence
;; FEATURE:
;; OTHER INFORMATION: Description of Artificial Sequence: Bridging
;; OTHER INFORMATION: oligonucleotides
US-09-626-528-3

Query Match 65.68; Score 11.8; DB 4; Length 40;
Best Local Similarity 60.08; Pred. No. 4.6e+02;
Matches 9; Conservative 4; Mismatches 2; Indels 0; Gaps 0;

QY 1 AAUGGCCUAUCGGUG 15
| :||| :|:|:|:|
Db 6 ACTGGCGTATCGGTG 20

RESULT 9
US-08-054-480-5
;; Sequence 5, Application US/08054480
;; Patent No. 5525504
;; GENERAL INFORMATION:
;; APPLICANT: Goebel, Werner
;; APPLICANT: Libby, Stephen
;; APPLICANT: Heffron, Fred
;; TITLE OF INVENTION: CYTOLYSIN GENE AND GENE PRODUCT
;; NUMBER OF SEQUENCES: 5
;; CORRESPONDENCE ADDRESS:
;; ADDRESSEE: MILLEN, WHITE, ZELANO, & BRANIGAN, P.C.
;; STREET: 2200 CLARENDON BOULEVARD, SUITE 1400
;; CITY: ARLINGTON
;; STATE: VIRGINIA
;; COUNTRY: USA
;; ZIP: 22201

;; COMPUTER READABLE FORM:
;; MEDIUM TYPE: Floppy disk
;; COMPUTER: IBM PC compatible
;; OPERATING SYSTEM: PC-DOS/MS-DOS
;; SOFTWARE: PatentIn Release #1.0, Version #1.25
;; CURRENT APPLICATION DATA:
;; APPLICATION NUMBER: US/08/054,480
;; FILING DATE: 04-APR-1993
;; CLASSIFICATION: 435
;; ATTORNEY/AGENT INFORMATION:
;; NAME: Heaney, Brian P.
;; REGISTRATION NUMBER: 32,542
;; REFERENCE/DOCKET NUMBER: MERCK 1496
;; TELECOMMUNICATION INFORMATION:
;; TELEPHONE: 703 243 6333
;; TELEFAX: 703 243 6410
;; TELEX: 64191
;; INFORMATION FOR SEQ ID NO: 5:
;; SEQUENCE CHARACTERISTICS:
;; LENGTH: 26 base pairs
;; TYPE: nucleic acid
;; STRANDEDNESS: single
;; TOPOLOGY: linear
;; MOLECULE TYPE: DNA (genomic)
;; ORIGINAL SOURCE:
;; ORGANISM: SALMONELLA
US-08-054-480-5

Query Match 64.4%; Score 11.6; DB 1; Length 26;
Best Local Similarity 66.7%; Pred. No. 5.7e+02;
Matches 12; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

QY 1 AAUGGCCUAUCGGUGCGA 18
|:|:|:| | |:|:|:|
Db 7 AATGGCAGAGAGGTGGCA 24

RESULT 10
US-08-413-803-10/c
;; Sequence 10, Application US/08413803
;; Patent No. 5766581
;; GENERAL INFORMATION:
;; APPLICANT: Bartley, Timothy D.
;; APPLICANT: Bogenberger, Jakob M.
;; APPLICANT: Bosselman, Robert A.
;; APPLICANT: Hunt, Pamela
;; APPLICANT: Kinstler, Olaf B.
;; APPLICANT: Samal, Babru B.
;; TITLE OF INVENTION: METHODS FOR TREATING MAMMALS WITH
;; TITLE OF INVENTION: MONO-PEGYLATED PROTEINS THAT STIMULATE MEGAKARYOCYTE
;; TITLE OF INVENTION: GROWTH AND DIFFERENTIATION
;; NUMBER OF SEQUENCES: 34
;; CORRESPONDENCE ADDRESS:
;; ADDRESSEE: AMGEN INC.
;; STREET: 1840 DeHavilland Drive
;; CITY: Thousand Oaks
;; STATE: California
;; COUNTRY: US
;; ZIP: 91320-1789

;; COMPUTER READABLE FORM:
;; MEDIUM TYPE: Floppy disk
;; COMPUTER: IBM PC compatible
;; OPERATING SYSTEM: PC-DOS/MS-DOS
;; SOFTWARE: PatentIn Release #1.0, Version #1.30
;; CURRENT APPLICATION DATA:
;; APPLICATION NUMBER: US/08/413,803
;; FILING DATE: 30-MAR-1995
;; CLASSIFICATION: 435
;; PRIOR APPLICATION DATA:
;; APPLICATION NUMBER: US 08/221,768
;; FILING DATE: 31-MAR-1994
;; PRIOR APPLICATION DATA:
;; APPLICATION NUMBER: US 08/252,628
;; FILING DATE: 31-MAY-1994
;; PRIOR APPLICATION DATA:
;; APPLICATION NUMBER: US 08/321,488
;; FILING DATE: 12-OCT-1994
;; PRIOR APPLICATION DATA:
;; APPLICATION NUMBER: US 08/347,780
;; FILING DATE: 30-NOV-1994
;; ATTORNEY/AGENT INFORMATION:
;; NAME: Cook Ph.D., Robert R.
;; REGISTRATION NUMBER: 31,602
;; REFERENCE/DOCKET NUMBER: A-290D
;; INFORMATION FOR SEQ ID NO: 10:
;; SEQUENCE CHARACTERISTICS:
;; LENGTH: 29 base pairs
;; TYPE: nucleic acid
;; STRANDEDNESS: single
;; TOPOLOGY: linear
;; MOLECULE TYPE: CDNA
US-08-413-803-10

Query Match 64.4%; Score 11.6; DB 1; Length 29;
Best Local Similarity 66.7%; Pred. No. 5.8e+02;
Matches 12; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

QY 1 AAUGGCCUAUCGGUGCGA 18
|:|:|:| | |:|:|:|
Db 19 AAAGGCGTATCCGGCGCA 2

RESULT 11
US-08-321-488A-10/c
;; Sequence 10, Application US/08321488A

```
; Patent No. 5795569
; GENERAL INFORMATION:
; APPLICANT: Bartley, Timothy D.
; APPLICANT: Bogenberger, Jakob M.
; APPLICANT: Bosselman, Robert A.
; APPLICANT: Hunt, Pamela
; APPLICANT: Kinstler, Olaf B.
; APPLICANT: Samal, Babru B.
; TITLE OF INVENTION: MONO-PEGYLATED PROTEINS THAT STIMULATE
; TITLE OF INVENTION: MEGAKARYOCYTE GROWTH AND DIFFERENTIATION
; NUMBER OF SEQUENCES: 27
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: AMGEN INC.
; STREET: 1840 DeHavilland Drive
; CITY: Thousand Oaks
; STATE: California
; COUNTRY: US
; ZIP: 91320-1789
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/321,488A
; FILING DATE: 12-OCT-1994
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/252,628
; FILING DATE: 31-MAY-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/221,768
; FILING DATE: 31-MAR-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Cook, Robert R.
; REGISTRATION NUMBER: 31,602
; REFERENCE/DOCKET NUMBER: A-290B
; INFORMATION FOR SEQ ID NO: 10:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 29 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: cDNA
; US-08-321-488A-10

Query Match 64.4%; Score 11.6; DB 1; Length 29;
Best Local Similarity 66.7%; Pred. No. 5.8e+02;
Matches 12; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

QY 1 AAUGGCCUAUCGGUGCGA 18
Db 19 AAAGGCGCTATCCGGCCGA 2

RESULT 12
PCT-US95-03776-10/c
; Sequence 10, Application PC/TUS9503776
; GENERAL INFORMATION:
; APPLICANT: AMGEN INC.
; TITLE OF INVENTION: Compositions and Methods for Stimulating
; TITLE OF INVENTION: Megakaryocyte Growth and Differentiation
; NUMBER OF SEQUENCES: 34
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Amgen Inc.
; STREET: 1840 Dehavilland Drive
; CITY: Thousand Oaks
; STATE: California
; COUNTRY: USA
; ZIP: 91320-1789
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
```

```
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: PCT/US95/03776
; FILING DATE:
; CLASSIFICATION:
; ATTORNEY/AGENT INFORMATION:
; NAME: Cook, Robert R.
; REFERENCE/DOCKET NUMBER: A-290-C
; INFORMATION FOR SEQ ID NO: 10:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 29 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: cDNA
; PCT-US95-03776-10

Query Match 64.4%; Score 11.6; DB 5; Length 29;
Best Local Similarity 66.7%; Pred. No. 5.8e+02;
Matches 12; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

QY 1 AAUGGCCUAUCGGUGCGA 18
Db 19 AAAGGCGCTATCCGGCCGA 2

RESULT 13
US-08-413-803-9/c
; Sequence 9, Application US/08413803
; Patent No. 5766581
; GENERAL INFORMATION:
; APPLICANT: Bartley, Timothy D.
; APPLICANT: Bogenberger, Jakob M.
; APPLICANT: Bosselman, Robert A.
; APPLICANT: Hunt, Pamela
; APPLICANT: Kinstler, Olaf B.
; APPLICANT: Samal, Babru B.
; TITLE OF INVENTION: METHODS FOR TREATING MAMMALS WITH
; TITLE OF INVENTION: MONO-PEGYLATED PROTEINS THAT STIMULATE MEGAKARYOCYTE
; GROWTH AND DIFFERENTIATION
; NUMBER OF SEQUENCES: 34
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: AMGEN INC.
; STREET: 1840 DeHavilland Drive
; CITY: Thousand Oaks
; STATE: California
; COUNTRY: US
; ZIP: 91320-1789
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/413,803
; FILING DATE: 30-MAR-1995
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/221,768
; FILING DATE: 31-MAR-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/252,628
; FILING DATE: 31-MAY-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/321,488
; FILING DATE: 12-OCT-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/347,780
; FILING DATE: 30-NOV-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Cook Ph.D., Robert R.
; REGISTRATION NUMBER: 31,602
```

REFERENCE/DOCKET NUMBER: A-290D
; INFORMATION FOR SEQ ID NO: 9:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 30 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: cDNA
US-08-413-803-9

Query Match 64.4%; Score 11.6; DB 1; Length 30;
Best Local Similarity 66.7%; Pred. No. 5.8e+02;
Matches 12; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

Qy 1 AAUGGCCUAUCGGUGCGA 18
|| ||||:|:| | |||
Db 19 AAAGGCCTATCCGGCGCA 2

RESULT 14
US-08-321-488A-9/c
; Sequence 9, Application US/08321488A
; Patent No. 5795569
; GENERAL INFORMATION:
; APPLICANT: Bartley, Timothy D.
; APPLICANT: Bogenberger, Jakob M.
; APPLICANT: Bosselman, Robert A.
; APPLICANT: Hunt, Pamela
; APPLICANT: Kinstler, Olaf B.
; APPLICANT: Samal, Babru B.
; TITLE OF INVENTION: MEGA-PGYLATED PROTEINS THAT STIMULATE
; TITLE OF INVENTION: MEGA-KARYOCYTE GROWTH AND DIFFERENTIATION
; NUMBER OF SEQUENCES: 27
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: AMGEN INC.
; STREET: 1840 DeHavilland Drive
; CITY: Thousand Oaks
; STATE: California
; COUNTRY: US
; ZIP: 91320-1789
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/321.488A
; FILING DATE: 12-OCT-1994
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/252,628
; FILING DATE: 31-MAY-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/221,768
; FILING DATE: 31-MAR-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Cook, Robert R.
; REGISTRATION NUMBER: 31,602
; REFERENCE/DOCKET NUMBER: A-290B
; INFORMATION FOR SEQ ID NO: 9:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 30 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: cDNA
US-08-321-488A-9

Query Match 64.4%; Score 11.6; DB 1; Length 30;
Best Local Similarity 66.7%; Pred. No. 5.8e+02;
Matches 12; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

Qy 1 AAUGGCCUAUCGGUGCGA 18

Db 19 AAAGGCCTATCCGGCGCA 2
|| ||||:|:| | |||

RESULT 15
US-08-943-915-14/c
; Sequence 14, Application US/08943915
; Patent No. 5998170
; GENERAL INFORMATION:
; APPLICANT: Itoh, No. 5998170uyuki
; APPLICANT: Martin, Frank
; APPLICANT: Danilenko, Dmitry
; TITLE OF INVENTION: A FIBROBLAST GROWTH FACTOR
; NUMBER OF SEQUENCES: 33
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Amgen Inc.
; STREET: 1840 DeHavilland Drive
; CITY: Thousand Oaks
; STATE: California
; COUNTRY: USA
; ZIP: 91320-1789
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/943,915
; FILING DATE:
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Mazza, Richard J.
; REGISTRATION NUMBER: 27,657
; REFERENCE/DOCKET NUMBER: A-469
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 805.447.4112
; TELEFAX: 805.447.1090
; INFORMATION FOR SEQ ID NO: 14:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 30 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: other nucleic acid
; DESCRIPTION: /desc = "OLIGONUCLEOTIDE"
US-08-943-915-14

Query Match 64.4%; Score 11.6; DB 2; Length 30;
Best Local Similarity 66.7%; Pred. No. 5.8e+02;
Matches 12; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

Qy 1 AAUGGCCUAUCGGUGCGA 18
|| ||||:|:| | |||
Db 19 AAAGGCCTATCCGGCGCA 2

Search completed: July 6, 2003, 15:30:04
Job time : 48.4545 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: July 6, 2003, 15:04:31 ; Search time 102 Seconds
(without alignments)
275.469 Million cell updates/sec

Title: US-09-780-929-98
Perfect score: 18
Sequence: 1 aaugccuauccgugcga 18

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 1085931 seqs, 780495707 residues

Total number of hits satisfying chosen parameters: 805896

Minimum DB seq length: 0
Maximum DB seq length: 60

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : Published_Applications_NA.*

- 1: /cgn2_6/ptodata/2/pubpna/US07_PUBCOMB.seq.*
- 2: /cgn2_6/ptodata/2/pubpna/PCT_NEW_PUB.seq.*
- 3: /cgn2_6/ptodata/2/pubpna/US06_NEW_PUB.seq.*
- 4: /cgn2_6/ptodata/2/pubpna/US06_PUBCOMB.seq.*
- 5: /cgn2_6/ptodata/2/pubpna/US07_NEW_PUB.seq.*
- 6: /cgn2_6/ptodata/2/pubpna/PCTUS_PUBCOMB.seq.*
- 7: /cgn2_6/ptodata/2/pubpna/US08_NEW_PUB.seq.*
- 8: /cgn2_6/ptodata/2/pubpna/US08_PUBCOMB.seq.*
- 9: /cgn2_6/ptodata/2/pubpna/US09_NEW_PUB.seq.*
- 10: /cgn2_6/ptodata/2/pubpna/US09_PUBCOMB.seq.*
- 11: /cgn2_6/ptodata/2/pubpna/US10_NEW_PUB.seq.*
- 12: /cgn2_6/ptodata/2/pubpna/US10_PUBCOMB.seq.*
- 13: /cgn2_6/ptodata/2/pubpna/US60_NEW_PUB.seq.*
- 14: /cgn2_6/ptodata/2/pubpna/US60_PUBCOMB.seq.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	ID	Description
1	18	100.0	18	10	US-09-780-929-98
2	18	100.0	29	10	US-09-780-929-107
3	13.8	76.7	30	10	US-09-798-675-7
4	13.8	76.7	30	10	US-09-798-675-8
5	12.2	67.8	29	9	US-09-975-719-144
6	11.8	65.6	24	9	US-09-940-185-3899
7	11.6	64.4	30	10	US-09-005-243-36
8	11.6	64.4	30	10	US-09-224-683-36
9	11.6	64.4	46	9	US-09-880-508-15
10	11.6	64.4	46	9	US-10-158-314-15
11	11.6	64.4	50	9	US-09-741-179A-10
12	11.6	64.4	50	9	US-09-741-179A-14
13	11.4	63.3	24	9	US-09-940-185-3596
14	11.4	63.3	41	10	US-09-759-272B-4
15	11.4	63.3	42	10	US-09-838-386-14
16	11.4	63.3	45	10	US-09-838-386-13
17	11.4	63.3	45	10	US-09-838-386-17
18	11.4	63.3	45	10	US-09-838-386-18
19	11.4	63.3	51	9	US-10-211-088-46

c	20	11.2	62.2	17	9	US-09-780-533A-633	Sequence 633, Appl
c	21	11.2	62.2	19	9	US-09-796-081-3	Sequence 3, Appli
c	22	11.2	62.2	19	9	US-09-796-081-4	Sequence 4, Appli
c	23	11.2	62.2	22	10	US-09-997-664-85	Sequence 85, Appl
c	24	11.2	62.2	25	9	US-10-098-263B-3120	Sequence 3120, Ap
c	25	11.2	62.2	25	9	US-10-098-263B-12236	Sequence 12236, A
c	26	11.2	62.2	25	9	US-10-098-263B-26400	Sequence 26400, A
c	27	11.2	62.2	25	9	US-10-098-263B-100906	Sequence 100906,
c	28	11.2	62.2	25	9	US-10-098-263B-120820	Sequence 120820,
c	29	11.2	62.2	25	9	US-10-098-263B-126689	Sequence 126689,
c	30	11.2	62.2	29	9	US-10-062-458-23	Sequence 23, Appl
c	31	11.2	62.2	30	12	US-10-053-632-11	Sequence 11, Appl
c	32	11.2	62.2	30	12	US-10-052-417-11	Sequence 11, Appl
c	33	11.2	62.2	40	10	US-09-245-802-75	Sequence 75, Appl
c	34	11.2	62.2	43	10	US-09-728-574-1	Sequence 1, Appli
c	35	11.2	62.2	44	9	US-10-062-458-18	Sequence 18, Appl
c	36	11.2	62.2	50	9	US-09-741-179A-13	Sequence 13, Appl
c	37	11.2	62.2	50	9	US-09-741-179A-15	Sequence 15, Appl
c	38	11.1	61.1	25	9	US-10-098-263B-26098	Sequence 26098, A
c	39	11.1	61.1	32	10	US-09-919-831-4	Sequence 4, Appli
c	40	10.8	60.0	21	9	US-09-995-529-223	Sequence 223, App
c	41	10.8	60.0	22	10	US-09-815-656-61	Sequence 61, Appl
c	42	10.8	60.0	22	10	US-09-815-656-63	Sequence 63, Appl
c	43	10.8	60.0	22	10	US-09-529-063-114	Sequence 114, App
c	44	10.8	60.0	24	9	US-09-940-185-1700	Sequence 1700, Ap
c	45	10.8	60.0	24	9	US-09-940-185-3740	Sequence 3740, Ap

ALIGNMENTS

RESULT 1
US-09-780-929-98
; Sequence 98, Application US/09780929
; Patent No. US20020151693A1
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc
; APPLICANT: Breaker, Ronald
; APPLICANT: Beigelman, Leo
; TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity
; FILE REFERENCE: MBH00-884-H (500/001)
; CURRENT APPLICATION NUMBER: US/09/780,929
; CURRENT FILING DATE: 2001-09-08
; PRIOR APPLICATION NUMBER: US 60/181,360
; PRIOR FILING DATE: 2000-02-08
; NUMBER OF SEQ ID NOS: 126
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 98
; LENGTH: 18
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid

US-09-780-929-98
Query Match 100.0%; Score 18; DB 10; Length 18;
Best Local Similarity 100.0%; Pred. No. 1.1;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AAUGGCCUUAUCGUGCGA 18
|||||
DB 1 AAUGGCCUUAUCGUGCGA 18

RESULT 2
US-09-780-929-107
; Sequence 107, Application US/09780929
; Patent No. US20020151693A1
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc
; APPLICANT: Breaker, Ronald
; APPLICANT: Beigelman, Leo
; TITLE OF INVENTION: Nucleic Acid Catalysts with Endonuclease Activity

```

; APPLICANT: Emory University
; TITLE OF INVENTION: HIV VACCINES
; FILE REFERENCE: E056 2020
; CURRENT APPLICATION NUMBER: US/09/798,675
; CURRENT FILING DATE: 2001-12-11
; PRIOR APPLICATION NUMBER: US 60/186,364
; PRIOR FILING DATE: 2000-03-02
; PRIOR APPLICATION NUMBER: US 60/251,083
; PRIOR FILING DATE: 2000-12-01
; NUMBER OF SEQ ID NOS: 24
; SOFTWARE: Patent version 3.0

```


US-09-940-185-3899

Query Match 65.6%; Score 11.8; DB 9; Length 24;
Best Local Similarity 73.3%; Pred. NO. 3.3e+03;
Matches 11; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 4 GGCCUACUGGCGCA 18
||||| ||:||||
DB 2 GGCTAGAGTGCGA 16

RESULT 7

US-09-005-243-36/c
; Sequence 36, Application US/09005243
; Patent No. US20020018763A1
; GENERAL INFORMATION:
; APPLICANT: Zsebo, Krisztina M.
; APPLICANT: Bosselman, Robert A.
; APPLICANT: Suggs, Sidney V.
; APPLICANT: Martin, Francis H.
; TITLE OF INVENTION: Stem Cell Factor
; NUMBER OF SEQUENCES: 104
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Marshall, O'Toole, Gerstein, Murray & Borun
; STREET: 6300 Sears Tower, 233 South Wacker Drive
; CITY: Chicago
; STATE: Illinois
; COUNTRY: United States of America
; ZIP: 60606-6402
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/005,243
; FILING DATE:
; CLASSIFICATION:

PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/449,653
FILING DATE: 24-MAY-1995
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/982,255
FILING DATE: 25-NOV-1992
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/589,701
FILING DATE: 01-OCT-1990
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/573,616
FILING DATE: 24-AUG-1990
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/537,198
FILING DATE: 11-JUN-1990
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/422,383
FILING DATE: 16-OCT-1989
ATTORNEY/AGENT INFORMATION:
NAME: Clough, David W.

REGISTRATION NUMBER: 36,107
REFERENCE/DOCKET NUMBER: 01017/34465
TELEPHONE: 312/474-6300
TELEFAX: 312/474-0448
TELEX: 25-3856

INFORMATION FOR SEQ ID NO: 36:
SEQUENCE CHARACTERISTICS:
LENGTH: 30 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA

US-09-005-243-36

Query Match 64.4%; Score 11.6; DB 10; Length 30;
Best Local Similarity 66.7%; Pred. NO. 4.3e+03;
Matches 12; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

QY 1 AAUGGCCUACUGGCGCA 18
|||||:|:| |||
DB 19 AAAGGCCTATCCGGCGA 2

RESULT 8

US-09-224-683-36/c
; Sequence 36, Application US/09224683
; Patent No. US20020031491A1
; GENERAL INFORMATION:
; APPLICANT: Zsebo, Krisztina M.
; APPLICANT: Bosselman, Robert A.
; APPLICANT: Suggs, Sidney V.
; APPLICANT: Martin, Francis H.
; TITLE OF INVENTION: Stem Cell Factor: Composition Claims
; NUMBER OF SEQUENCES: 104
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Marshall, O'Toole, Gerstein, Murray & Borun
; STREET: 6300 Sears Tower, 233 South Wacker Drive
; CITY: Chicago
; STATE: Illinois
; COUNTRY: United States of America
; ZIP: 60606-6402
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/224,683
; FILING DATE:
; CLASSIFICATION:

PRIOR APPLICATION DATA:
APPLICATION NUMBER: 09/005,893
FILING DATE: 12-JAN-1998
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/449,653
FILING DATE: 24-MAY-1995
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/982,255
FILING DATE: 25-NOV-1992
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/589,701
FILING DATE: 01-OCT-1990
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/573,616
FILING DATE: 24-AUG-1990
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/537,198
FILING DATE: 11-JUN-1990
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/422,383
FILING DATE: 16-OCT-1989
ATTORNEY/AGENT INFORMATION:
NAME: Clough, David W.

REGISTRATION NUMBER: 36,107
REFERENCE/DOCKET NUMBER: 01017/35136
TELEPHONE: 312/474-6300
TELEFAX: 312/474-0448
TELEX: 25-3856

INFORMATION FOR SEQ ID NO: 36:
SEQUENCE CHARACTERISTICS:
LENGTH: 30 base pairs
TYPE: nucleic acid
STRANDEDNESS: single

Oy 1 AAUGGCCUAUCGG 13
||:|||||:|
Db 23 AATGCCCTATTGG 35

Search completed: July 6, 2003, 16:52:33
Job time : 103 secs

GenCore version 5.1.1.6
Copyright (c) 1993 - 2003 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: July 6, 2003, 14:40:47 ; Search time 1951.09 Seconds
(without alignments)
231.955 Million cell updates/sec

Title: US-09-780-929-98
Perfect score: 18
Sequence: 1 aauggcuaucgugcgga 18
Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 24791104 seqs, 12571243825 residues

Total number of hits satisfying chosen parameters: 12745074

Minimum DB seq length: 0
Maximum DB seq length: 60

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :

Pending Patents_NA_Main.*
1: /cgn2_6/ptodata/1/pna/PCTUS_COMB.seq.*
2: /cgn2_6/ptodata/1/pna/US06_COMB.seq.*
3: /cgn2_6/ptodata/1/pna/US07_COMB.seq.*
4: /cgn2_6/ptodata/1/pna/US08_COMB.seq.*
5: /cgn2_6/ptodata/1/pna/US081_COMB.seq.*
6: /cgn2_6/ptodata/1/pna/US082_COMB.seq.*
7: /cgn2_6/ptodata/1/pna/US083_COMB.seq.*
8: /cgn2_6/ptodata/1/pna/US084_COMB.seq.*
9: /cgn2_6/ptodata/1/pna/US085_COMB.seq.*
10: /cgn2_6/ptodata/1/pna/US086_COMB.seq.*
11: /cgn2_6/ptodata/1/pna/US087_COMB.seq.*
12: /cgn2_6/ptodata/1/pna/US088_COMB.seq.*
13: /cgn2_6/ptodata/1/pna/US089_COMB.seq.*
14: /cgn2_6/ptodata/1/pna/US090_COMB.seq.*
15: /cgn2_6/ptodata/1/pna/US091_COMB.seq.*
16: /cgn2_6/ptodata/1/pna/US092_COMB.seq.*
17: /cgn2_6/ptodata/1/pna/US093_COMB.seq.*
18: /cgn2_6/ptodata/1/pna/US094_COMB.seq.*
19: /cgn2_6/ptodata/1/pna/US095A_COMB.seq.*
20: /cgn2_6/ptodata/1/pna/US095B_COMB.seq.*
21: /cgn2_6/ptodata/1/pna/US095C_COMB.seq.*
22: /cgn2_6/ptodata/1/pna/US095D_COMB.seq.*
23: /cgn2_6/ptodata/1/pna/US096A_COMB.seq.*
24: /cgn2_6/ptodata/1/pna/US096B_COMB.seq.*
25: /cgn2_6/ptodata/1/pna/US096C_COMB.seq.*
26: /cgn2_6/ptodata/1/pna/US096D_COMB.seq.*
27: /cgn2_6/ptodata/1/pna/US096E_COMB.seq.*
28: /cgn2_6/ptodata/1/pna/US097A_COMB.seq.*
29: /cgn2_6/ptodata/1/pna/US097B_COMB.seq.*
30: /cgn2_6/ptodata/1/pna/US097C_COMB.seq.*
31: /cgn2_6/ptodata/1/pna/US098A_COMB.seq.*
32: /cgn2_6/ptodata/1/pna/US098B_COMB.seq.*
33: /cgn2_6/ptodata/1/pna/US098C_COMB.seq.*
34: /cgn2_6/ptodata/1/pna/US099A_COMB.seq.*
35: /cgn2_6/ptodata/1/pna/US099B_COMB.seq.*
36: /cgn2_6/ptodata/1/pna/US099C_COMB.seq.*
37: /cgn2_6/ptodata/1/pna/US099D_COMB.seq.*
38: /cgn2_6/ptodata/1/pna/US100A_COMB.seq.*
39: /cgn2_6/ptodata/1/pna/US100B_COMB.seq.*
40: /cgn2_6/ptodata/1/pna/US101A_COMB.seq.*
41: /cgn2_6/ptodata/1/pna/US101B_COMB.seq.*
42: /cgn2_6/ptodata/1/pna/US102A_COMB.seq.*
43: /cgn2_6/ptodata/1/pna/US102B_COMB.seq.*

44: /cgn2_6/ptodata/1/pna/US6000_COMB.seq.*
45: /cgn2_6/ptodata/1/pna/US6001_COMB.seq.*
46: /cgn2_6/ptodata/1/pna/US6002_COMB.seq.*
47: /cgn2_6/ptodata/1/pna/US6003_COMB.seq.*
48: /cgn2_6/ptodata/1/pna/US6004_COMB.seq.*
49: /cgn2_6/ptodata/1/pna/US6005_COMB.seq.*
50: /cgn2_6/ptodata/1/pna/US6006_COMB.seq.*
51: /cgn2_6/ptodata/1/pna/US6007_COMB.seq.*
52: /cgn2_6/ptodata/1/pna/US6008_COMB.seq.*
53: /cgn2_6/ptodata/1/pna/US6009_COMB.seq.*
54: /cgn2_6/ptodata/1/pna/US6010_COMB.seq.*
55: /cgn2_6/ptodata/1/pna/US6011_COMB.seq.*
56: /cgn2_6/ptodata/1/pna/US6012_COMB.seq.*
57: /cgn2_6/ptodata/1/pna/US6013_COMB.seq.*
58: /cgn2_6/ptodata/1/pna/US6014_COMB.seq.*
59: /cgn2_6/ptodata/1/pna/US6015_COMB.seq.*
60: /cgn2_6/ptodata/1/pna/US6016_COMB.seq.*
61: /cgn2_6/ptodata/1/pna/US6017_COMB.seq.*
62: /cgn2_6/ptodata/1/pna/US6018_COMB.seq.*
63: /cgn2_6/ptodata/1/pna/US6019_COMB.seq.*
64: /cgn2_6/ptodata/1/pna/US6020_COMB.seq.*
65: /cgn2_6/ptodata/1/pna/US6021_COMB.seq.*
66: /cgn2_6/ptodata/1/pna/US6022_COMB.seq.*
67: /cgn2_6/ptodata/1/pna/US6023_COMB.seq.*
68: /cgn2_6/ptodata/1/pna/US6024_COMB.seq.*
69: /cgn2_6/ptodata/1/pna/US6025_COMB.seq.*
70: /cgn2_6/ptodata/1/pna/US6026_COMB.seq.*
71: /cgn2_6/ptodata/1/pna/US6027_COMB.seq.*
72: /cgn2_6/ptodata/1/pna/US6028_COMB.seq.*
73: /cgn2_6/ptodata/1/pna/US6029_COMB.seq.*
74: /cgn2_6/ptodata/1/pna/US6030_COMB.seq.*
75: /cgn2_6/ptodata/1/pna/US6031_COMB.seq.*
76: /cgn2_6/ptodata/1/pna/US6032_COMB.seq.*
77: /cgn2_6/ptodata/1/pna/US6033_COMB.seq.*
78: /cgn2_6/ptodata/1/pna/US6034_COMB.seq.*
79: /cgn2_6/ptodata/1/pna/US6035_COMB.seq.*
80: /cgn2_6/ptodata/1/pna/US6036_COMB.seq.*
81: /cgn2_6/ptodata/1/pna/US6037_COMB.seq.*
82: /cgn2_6/ptodata/1/pna/US6038_COMB.seq.*
83: /cgn2_6/ptodata/1/pna/US6039_COMB.seq.*
84: /cgn2_6/ptodata/1/pna/US6040_COMB.seq.*
85: /cgn2_6/ptodata/1/pna/US6041_COMB.seq.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	18	100.0	18	30	US-09-780-929-98
2	18	100.0	29	30	US-09-780-929-107
c 3	13.8	76.7	30	1	PCT-US01-06795A-7
c 5	13.8	76.7	30	1	PCT-US01-06795A-8
c 6	13.8	76.7	30	30	US-09-798-675-7
7	13.4	74.4	30	30	US-09-798-675-8
8	13.4	74.4	60	34	US-09-908-975-20491
9	13.4	74.4	60	34	US-09-908-975A-20491
10	13.2	73.3	25	36	US-09-954-427-30098
c 11	13.2	73.3	25	36	US-09-956-584-426597
c 12	13.2	73.3	25	36	US-09-956-604-6465
c 13	13.2	73.3	25	36	US-09-956-604-105593
c 14	13.2	73.3	25	36	US-09-956-604-114271
c 15	13.2	73.3	25	36	US-09-956-604A-6465
c 16	13.2	73.3	25	36	US-09-956-604A-105593
c 17	13.2	73.3	25	36	US-09-956-604A-114271
c 18	13.2	73.3	25	36	US-09-956-604B-6465
c 19	13.2	73.3	25	36	US-09-956-604B-105593
c 20	13.2	73.3	25	36	US-09-956-604B-114271
21	13.2	73.3	25	67	US-60-233-166-30098
					Sequence 98, Appl
					Sequence 107, Appl
					Sequence 7, Appl
					Sequence 8, Appl
					Sequence 7, Appl
					Sequence 8, Appl
					Sequence 20491, A
					Sequence 20491, A
					Sequence 20491, A
					Sequence 30098, A
					Sequence 426597, A
					Sequence 6465, Ap
					Sequence 105593, A
					Sequence 114271, A
					Sequence 105593, A
					Sequence 105593, A
					Sequence 114271, A
					Sequence 6465, Ap
					Sequence 105593, A
					Sequence 114271, A
					Sequence 30098, A

PCT-US01-06795A-8

Query Match 76.7%; Score 13.8; DB 1; Length 30;
Best Local Similarity 64.7%; Pred. No. 2.4e+03;
Matches 11; Conservative 4; Mismatches 2; Indels 0; Gaps 0;

Qy 2 AUGGCCUACGUGCGGA 18
1:1111:1:11:1111
Db 6 ATGGCGTATCGATCGGA 22

RESULT 5

US-09-798-675-7/c
; Sequence 7, Application US/09798675
; GENERAL INFORMATION:
; APPLICANT: Emory University
; TITLE OF INVENTION: HIV VACCINES
; FILE REFERENCE: E056 2020
; CURRENT APPLICATION NUMBER: US/09/798,675
; CURRENT FILING DATE: 2001-12-11
; PRIOR APPLICATION NUMBER: US 60/186,364
; PRIOR FILING DATE: 2000-03-02
; PRIOR APPLICATION NUMBER: US 60/251,083
; PRIOR FILING DATE: 2000-12-01
; NUMBER OF SEQ ID NOS: 24
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 7
; LENGTH: 30
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:

OTHER INFORMATION: primer for site-directed mutagenesis for introducing Cla I site

Query Match 76.7%; Score 13.8; DB 30; Length 30;
Best Local Similarity 64.7%; Pred. No. 2.4e+03;
Matches 11; Conservative 4; Mismatches 2; Indels 0; Gaps 0;

Qy 2 AUGGCCUACGUGCGGA 18
1:1111:1:11:1111
Db 25 ATGGCGTATCGATCGGA 9

RESULT 6

US-09-798-675-8
; Sequence 8, Application US/09798675
; GENERAL INFORMATION:
; APPLICANT: Emory University
; TITLE OF INVENTION: HIV VACCINES
; FILE REFERENCE: E056 2020
; CURRENT APPLICATION NUMBER: US/09/798,675
; CURRENT FILING DATE: 2001-12-11
; PRIOR APPLICATION NUMBER: US 60/186,364
; PRIOR FILING DATE: 2000-03-02
; PRIOR APPLICATION NUMBER: US 60/251,083
; PRIOR FILING DATE: 2000-12-01
; NUMBER OF SEQ ID NOS: 24
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 8
; LENGTH: 30
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:

OTHER INFORMATION: primer for site-directed mutagenesis to introduce Cla I site

Query Match 76.7%; Score 13.8; DB 30; Length 30;
Best Local Similarity 64.7%; Pred. No. 2.4e+03;
Matches 11; Conservative 4; Mismatches 2; Indels 0; Gaps 0;

Qy 2 AUGGCCUACGUGCGGA 18
1:1111:1:11:1111
Db 6 ATGGCGTATCGATCGGA 22

US-09-908-975-20491
; Sequence 20491, Application US/09908975
; GENERAL INFORMATION:
; APPLICANT: SHOSHAN, Avi
; APPLICANT: WASSERMAN, Alon
; APPLICANT: MINTZ, Eli
; APPLICANT: MINTZ, Liat
; APPLICANT: FAIGLER, Simchon
; TITLE OF INVENTION: OLIGONUCLEOTIDE LIBRARY FOR DETECTING RNA TRANSCRIPTS AND SI
; FILE REFERENCE: 36688-0005
; CURRENT APPLICATION NUMBER: US/09/908,975
; CURRENT FILING DATE: 2001-07-20
; PRIOR APPLICATION NUMBER: US 60/287,724
; PRIOR FILING DATE: 2001-05-02
; PRIOR APPLICATION NUMBER: US 60/221,607
; PRIOR FILING DATE: 2000-07-28
; NUMBER OF SEQ ID NOS: 32337
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 20491
; LENGTH: 60
; TYPE: DNA
; ORGANISM: Homo sapiens
; US-09-908-975-20491

Query Match 74.4%; Score 13.4; DB 34; Length 60;
Best Local Similarity 66.7%; Pred. No. 4.4e+03;
Matches 10; Conservative 4; Mismatches 1; Indels 0; Gaps 0;

Qy 1 AAUGGCCUACGUGG 15
1:1111:1:11:11
Db 1 ATTGGCCTATCGGTG 15

RESULT 8

US-09-908-975A-20491
; Sequence 20491, Application US/09908975A
; GENERAL INFORMATION:
; APPLICANT: SHOSHAN, Avi
; APPLICANT: WASSERMAN, Alon
; APPLICANT: MINTZ, Eli
; APPLICANT: MINTZ, Liat
; APPLICANT: FAIGLER, Simchon
; TITLE OF INVENTION: OLIGONUCLEOTIDE LIBRARY FOR DETECTING RNA TRANSCRIPTS AND SI
; FILE REFERENCE: 36688-0006
; CURRENT APPLICATION NUMBER: US/09/908,975A
; CURRENT FILING DATE: 2001-07-20
; PRIOR APPLICATION NUMBER: US 60/287,724
; PRIOR FILING DATE: 2001-05-02
; PRIOR APPLICATION NUMBER: US 60/221,607
; PRIOR FILING DATE: 2000-07-28
; NUMBER OF SEQ ID NOS: 32337
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 20491
; LENGTH: 60
; TYPE: DNA
; ORGANISM: Homo sapiens
; US-09-908-975A-20491

Query Match 74.4%; Score 13.4; DB 34; Length 60;
Best Local Similarity 66.7%; Pred. No. 4.4e+03;
Matches 10; Conservative 4; Mismatches 1; Indels 0; Gaps 0;

Qy 1 AAUGGCCUACGUGG 15
1:1111:1:11:11
Db 1 ATTGGCCTATCGGTG 15

RESULT 9

```
US-60-287-724-20491
; Sequence 20491, Application US/60287724
; GENERAL INFORMATION:
; APPLICANT: SHOSHAN, Avi
; APPLICANT: WASSERMAN, Alon
; APPLICANT: MINTZ, Eli
; APPLICANT: FAIGLER, Simchon
; TITLE OF INVENTION: OLIGONUCLEOTIDE LIBRARY FOR DETECTING RNA TRANSCRIPTS AND SPLICING
; TITLE OF INVENTION: THAT POPULATE A TRANSCRIPTOME
; FILE REFERENCE: 36688-0004
; CURRENT APPLICATION NUMBER: US/60/287,724
; CURRENT FILING DATE: 2001-05-02
; NUMBER OF SEQ ID NOS: 32337
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 20491
; LENGTH: 60
; TYPE: DNA
; ORGANISM: Homo sapiens
US-60-287-724-20491

Query Match      74.4%; Score 13.4; DB 72; Length 60;
Best Local Similarity 66.7%; Pred. No. 4.4e+03;
Matches 10; Conservative 4; Mismatches 1; Indels 0; Gaps 0;

Qy      1 AAUGGCCUAUCGGUG 15
       1 :||||:|:||||:|
Db      1 ATTGGCCTATCGGTG 15

RESULT 10
US-09-954-427-30098
; Sequence 30098, Application US/09954427
; GENERAL INFORMATION:
; APPLICANT: Mittmann
; APPLICANT: Affymetrix, Inc.
; TITLE OF INVENTION: Methods of Genetic Analysis of the Rat
; TITLE OF INVENTION: Genome
; FILE REFERENCE: 3112
; CURRENT APPLICATION NUMBER: US/09/954,427
; CURRENT FILING DATE: 2001-09-17
; NUMBER OF SEQ ID NOS: 420907
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 30098
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Rattus norvegicus
; PUBLICATION INFORMATION:
; DATABASE ACCESSION NUMBER: GenBank AA818650
US-09-954-427-30098

Query Match      73.3%; Score 13.2; DB 36; Length 25;
Best Local Similarity 66.7%; Pred. No. 5.3e+03;
Matches 12; Conservative 3; Mismatches 3; Indels 0; Gaps 0;

Qy      1 AAUGGCCUAUCGGUGCA 18
       1 :||||:|:||||:|
Db      8 AATGCTCTATCGCGCAA 25

RESULT 11
US-09-956-584-426597/c
; Sequence 426597, Application US/09956584
; GENERAL INFORMATION:
; APPLICANT: Mittmann, Michael
; TITLE OF INVENTION: Methods of Genetic Analysis of Mus Musculus
; FILE REFERENCE: 3115.1
; CURRENT APPLICATION NUMBER: US/09/956,584
; CURRENT FILING DATE: 2001-09-19
; PRIOR APPLICATION NUMBER: 60/234,017
; PRIOR FILING DATE: 2000-09-20
; NUMBER OF SEQ ID NOS: 605887
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1

US-09-956-604-426597
; SEQ ID NO 426597
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Mus musculus
US-09-956-584-426597

Query Match      73.3%; Score 13.2; DB 36; Length 25;
Best Local Similarity 66.7%; Pred. No. 5.3e+03;
Matches 12; Conservative 3; Mismatches 3; Indels 0; Gaps 0;

Qy      1 AAUGGCCUAUCGGUGCGCA 18
       1 :||||:|:||||:|
Db      21 AAGGACCTATCGGTCCGA 4

RESULT 12
US-09-956-604-6465/c
; Sequence 6465, Application US/09956604
; GENERAL INFORMATION:
; APPLICANT: Mittmann, Michael
; TITLE OF INVENTION: Methods of Genetic Analysis of Escherichia coli
; FILE REFERENCE: 3117.1
; CURRENT APPLICATION NUMBER: US/09/956,604
; CURRENT FILING DATE: 2001-09-19
; PRIOR APPLICATION NUMBER: 60/234,049
; PRIOR FILING DATE: 2000-09-19
; NUMBER OF SEQ ID NOS: 141629
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 6465
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Escherichia coli
US-09-956-604-6465

Query Match      73.3%; Score 13.2; DB 36; Length 25;
Best Local Similarity 61.1%; Pred. No. 5.3e+03;
Matches 11; Conservative 4; Mismatches 3; Indels 0; Gaps 0;

Qy      1 AAUGGCCUAUCGGUGCGCA 18
       1 :||||:|:||||:|
Db      23 AATGGTCTATCGTTGAGA 6

RESULT 13
US-09-956-604-105593/c
; Sequence 105593, Application US/09956604
; GENERAL INFORMATION:
; APPLICANT: Mittmann, Michael
; TITLE OF INVENTION: Methods of Genetic Analysis of Escherichia coli
; FILE REFERENCE: 3117.1
; CURRENT APPLICATION NUMBER: US/09/956,604
; CURRENT FILING DATE: 2001-09-19
; PRIOR APPLICATION NUMBER: 60/234,049
; PRIOR FILING DATE: 2000-09-19
; NUMBER OF SEQ ID NOS: 141629
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 105593
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Escherichia coli
US-09-956-604-105593

Query Match      73.3%; Score 13.2; DB 36; Length 25;
Best Local Similarity 61.1%; Pred. No. 5.3e+03;
Matches 11; Conservative 4; Mismatches 3; Indels 0; Gaps 0;

Qy      1 AAUGGCCUAUCGGUGCGCA 18
       1 :||||:|:||||:|
Db      20 AGTGGCCTATCAGTGCCA 3

RESULT 14
US-09-956-604-114271/c
```



```
; Sequence 114271, Application US/09956604
; GENERAL INFORMATION:
; APPLICANT: Mittmann, Michael
; TITLE OF INVENTION: Methods of Genetic Analysis of Escherichia coli
; FILE REFERENCE: 3117.1
; CURRENT APPLICATION NUMBER: US/09/956,604
; CURRENT FILING DATE: 2001-09-19
; PRIOR APPLICATION NUMBER: 60/234, 049
; PRIOR FILING DATE: 2000-09-19
; NUMBER OF SEQ ID NOS: 141629
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 114271
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Escherichia coli
US-09-956-604-114271

Query Match      73.3%; Score 13.2; DB 36; Length 25;
Best Local Similarity 61.1%; Pred. No. 5.3e+03;
Matches 11; Conservative 4; Mismatches 3; Indels 0; Gaps 0;
```

```
QY      1  AAUGGCCUAUCGUGCGGA 18
DB      20  AGTGGCTATCGCTACGA 3

RESULT 15
US-09-956-604A-6465/c
; Sequence 6465, Application US/09956604A
; GENERAL INFORMATION:
; APPLICANT: Mittmann, Michael
; TITLE OF INVENTION: Methods of Genetic Analysis of Escherichia coli
; FILE REFERENCE: 3117.1
; CURRENT APPLICATION NUMBER: US/09/956,604A
; CURRENT FILING DATE: 2001-09-19
; PRIOR APPLICATION NUMBER: 60/234, 049
; PRIOR FILING DATE: 2000-09-19
; NUMBER OF SEQ ID NOS: 141629
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 6465
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Escherichia coli
US-09-956-604A-6465
```

```
Query Match      73.3%; Score 13.2; DB 36; Length 25;
Best Local Similarity 61.1%; Pred. No. 5.3e+03;
Matches 11; Conservative 4; Mismatches 3; Indels 0; Gaps 0;
```

```
QY      1  AAUGGCCUAUCGUGCGGA 18
DB      23  AATGGTCTATCGTTGAGA 6

Search completed: July 6, 2003, 16:29:55
Job time : 1953.09 secs
```

GenCore version 5.1.6
Copyright (c) 1993 - 2003 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: July 6, 2003, 14:51:21 ; Search time 624.545 Seconds

(without alignments)
206.249 Million cell updates/sec

Title: US-09-780-929-98

Perfect score: 18

Sequence: 1 aaugccuauaggugcga 18

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 8255821 seqs, 3578102051 residues

Total number of hits satisfying chosen parameters: 9359164

Minimum DB seq length: 0

Maximum DB seq length: 60

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Pending_Patents_NA_New.*

- 1: /cgn2_6/ptodata/2/pna/PCT_NEW_COMB.seq.*
- 2: /cgn2_6/ptodata/2/pna/PCT_NEW_COMB.seq.*
- 3: /cgn2_6/ptodata/2/pna/US06_NEW_COMB.seq.*
- 4: /cgn2_6/ptodata/2/pna/US06_NEW_COMB.seq.*
- 5: /cgn2_6/ptodata/2/pna/US07_NEW_COMB.seq.*
- 6: /cgn2_6/ptodata/2/pna/US07_NEW_COMB.seq.*
- 7: /cgn2_6/ptodata/2/pna/US08_NEW_COMB.seq.*
- 8: /cgn2_6/ptodata/2/pna/US08_NEW_COMB.seq.*
- 9: /cgn2_6/ptodata/2/pna/US09_NEW_COMB.seq.*
- 10: /cgn2_6/ptodata/2/pna/US09_NEW_COMB.seq.*
- 11: /cgn2_6/ptodata/2/pna/US09_NEW_COMB.seq.*
- 12: /cgn2_6/ptodata/2/pna/US09_NEW_COMB.seq.*
- 13: /cgn2_6/ptodata/2/pna/US10_NEW_COMB.seq.*
- 14: /cgn2_6/ptodata/2/pna/US10_NEW_COMB.seq.*
- 15: /cgn2_6/ptodata/2/pna/US10_NEW_COMB.seq.*
- 16: /cgn2_6/ptodata/2/pna/US10_NEW_COMB.seq.*
- 17: /cgn2_6/ptodata/2/pna/US60_NEW_COMB.seq.*
- 18: /cgn2_6/ptodata/2/pna/US60_NEW_COMB.seq.*
- 19: /cgn2_6/ptodata/2/pna/US60_NEW_COMB.seq.*
- 20: /cgn2_6/ptodata/2/pna/US60_NEW_COMB.seq.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
C 1	13.8	76.7	30	15 US-10-336-566-21	Sequence 21, Appl
C 2	13.8	76.7	30	15 US-10-336-566-22	Sequence 22, Appl
C 3	13.2	73.3	24	14 US-10-310-188-56966	Sequence 56966, A
C 4	13.2	73.3	25	18 US-60-427-808-19526	Sequence 19526, A
C 5	13.2	73.3	25	18 US-60-427-808-125229	Sequence 125229, A
C 6	13.2	73.3	25	18 US-60-427-808-897906	Sequence 897906, A
C 7	13.2	73.3	25	20 US-60-475-871-64253	Sequence 64253, A
C 8	13.2	73.3	28	15 US-10-287-787-12927	Sequence 12927, A
C 9	13.2	73.3	43	15 US-10-287-787-21425	Sequence 21425, A
C 10	13.2	73.3	48	15 US-10-287-787-4996	Sequence 4996, Ap
C 11	13	72.2	25	14 US-10-355-577-821852	Sequence 821852, A
C 12	13	72.2	25	18 US-60-427-808-512279	Sequence 512279, A
C 13	12.8	71.1	25	11 US-09-954-445A-39955	Sequence 39955, A

14	12.8	71.1	25	14	US-10-355-577-133076	Sequence 133076,
C 15	12.8	71.1	25	14	US-10-355-577-152466	Sequence 152466,
C 16	12.8	71.1	25	14	US-10-355-577-703461	Sequence 703461,
C 17	12.8	71.1	25	14	US-10-355-577-834630	Sequence 834630,
C 18	12.8	71.1	25	14	US-10-355-577-904618	Sequence 904618,
C 19	12.8	71.1	25	18	US-60-427-808-56539	Sequence 56539, A
C 20	12.8	71.1	25	18	US-60-427-808-238519	Sequence 238519,
C 21	12.8	71.1	25	18	US-60-427-808-304674	Sequence 304674,
C 22	12.8	71.1	25	18	US-60-427-808-534678	Sequence 534678,
C 23	12.8	71.1	25	18	US-60-427-808-547438	Sequence 547438,
C 24	12.8	71.1	25	18	US-60-427-836-69606	Sequence 69606, A
C 25	12.8	71.1	25	18	US-60-427-836-81651	Sequence 81651, A
C 26	12.8	71.1	25	18	US-60-427-836-204586	Sequence 204586,
C 27	12.8	71.1	25	18	US-60-427-836-658226	Sequence 658226,
C 28	12.8	71.1	25	19	US-60-469-545-81455	Sequence 81455, A
C 29	12.8	71.1	25	19	US-60-469-545-81457	Sequence 81457, A
C 30	12.8	71.1	25	19	US-60-469-545-82435	Sequence 82435, A
C 31	12.8	71.1	25	19	US-60-469-545-155742	Sequence 155742,
C 32	12.8	71.1	25	19	US-60-469-545-156720	Sequence 156720,
C 33	12.8	71.1	25	19	US-60-469-545-156722	Sequence 156722,
C 34	12.4	68.9	25	12	US-09-660-0808-18293	Sequence 18293, A
C 35	12.4	68.9	25	14	US-10-355-577-565114	Sequence 565114,
C 36	12.4	68.9	25	18	US-60-427-808-918802	Sequence 918802,
C 37	12.4	68.9	25	18	US-60-427-836-63726	Sequence 63726, A
C 38	12.4	68.9	25	18	US-60-427-836-165832	Sequence 165832,
C 39	12.4	68.9	25	20	US-60-475-871-111969	Sequence 111969,
C 40	12.2	67.8	25	11	US-09-954-445A-10275	Sequence 10275, A
C 41	12.2	67.8	25	14	US-10-355-577-446254	Sequence 446254,
C 42	12.2	67.8	25	14	US-10-355-577-501905	Sequence 501905,
C 43	12.2	67.8	25	14	US-10-355-577-650463	Sequence 650463,
C 44	12.2	67.8	25	14	US-10-355-577-732834	Sequence 732834,
C 45	12.2	67.8	25	14	US-10-355-577-742769	Sequence 742769,

ALIGNMENTS

RESULT 1
US-10-336-566-21/c
; Sequence 21, Application US/10336566
; GENERAL INFORMATION:
; APPLICANT: Robinson, Harriet L.
; APPLICANT: Smith, James M.
; APPLICANT: Hua, Jian
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR GENERATING
; TITLE OF INVENTION: AN IMMUNE RESPONSE
; FILE REFERENCE: 12804-006001
; CURRENT APPLICATION NUMBER: US/10/336,566
; CURRENT FILING DATE: 2003-01-03
; PRIOR APPLICATION NUMBER: US 10/093,953
; PRIOR FILING DATE: 2002-03-08
; PRIOR APPLICATION NUMBER: US 09/798,675
; PRIOR FILING DATE: 2001-03-01
; PRIOR APPLICATION NUMBER: PCT/US01/06795
; PRIOR FILING DATE: 2001-03-02
; PRIOR APPLICATION NUMBER: US 60/251,083
; PRIOR FILING DATE: 2000-12-01
; PRIOR APPLICATION NUMBER: US 60/186,364
; PRIOR FILING DATE: 2000-03-02
; PRIOR APPLICATION NUMBER: US 60/324,845
; PRIOR FILING DATE: 2001-09-25
; NUMBER OF SEQ ID NOS: 87
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 21
; LENGTH: 30
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Primer
US-10-336-566-21
Query Match 76.7%; Score 13.8; DB 15; Length 30;
Best Local Similarity 64.7%; Pred. No. 9.6e+02;

FILE REFERENCE: JIM Zegeer Law Offices - 703-684-8333

```

Query Match      72.2%; Score 13; DB 14; Length 25;
Best Local Similarity 76.9%; Pred. No. 2.7e+03;
Matches 10; Conservative 3; Mismatches 0; Indels 0; Gaps 0;
ov      6 CCUAGCGGCGCA 18

```

```
Db          2 CCTATCGGTGCGA 14
          ||:|:|:|:|:|:|
RESULT 12
US-60-427-808-512279/c
; Sequence 512279, Application US/60427808
; GENERAL INFORMATION:
; APPLICANT: Xue Mei Zhou
; TITLE OF INVENTION: Methods of Genetic Analysis of Mouse
; FILE REFERENCE: 3528
; CURRENT APPLICATION NUMBER: US/60/427,808
; CURRENT FILING DATE: 2002-11-20
; NUMBER OF SEQ ID NOS: 982914
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 512279
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Mus musculus
US-60-427-808-512279
Query Match          72.2%; Score 13; DB 18; Length 25;
Best Local Similarity 76.8%; Pred. No. 2.7e+03;
Matches 10; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY          1 A AUGGCCCUAUCGG 13
          ||:|:|:|:|:|:|
Db          23 AATGGCCTATCGG 11

RESULT 13
US-09-954-445A-39955/c
; Sequence 39955, Application US/09954445A
; GENERAL INFORMATION:
; APPLICANT: Mittmann, Michael
; TITLE OF INVENTION: Methods of Genetic Analysis of Arabidopsis Thaliana
; FILE REFERENCE: 3116.1
; CURRENT APPLICATION NUMBER: US/09/954,445A
; CURRENT FILING DATE: 2000-09-17
; PRIOR APPLICATION NUMBER: 60/233,620
; PRIOR FILING DATE: 2000-09-18
; NUMBER OF SEQ ID NOS: 131820
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 39955
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Arabidopsis thaliana
US-09-954-445A-39955
Query Match          71.1%; Score 12.8; DB 11; Length 25;
Best Local Similarity 62.5%; Pred. No. 3.5e+03;
Matches 10; Conservative 4; Mismatches 2; Indels 0; Gaps 0;

QY          1 A AUGGCCCUAUCGGUCG 16
          ||:|:|:|:|:|:|
Db          22 AATGCATATCGGTGC 7

RESULT 14
US-10-355-577-133076
; Sequence 133076, Application US/10355577
; GENERAL INFORMATION:
; APPLICANT: Mittmann, Michael
; TITLE OF INVENTION: Methods of Genetic Analysis of Probes: HG-U133
; FILE REFERENCE: 3121
; CURRENT APPLICATION NUMBER: US/10/355,577
; CURRENT FILING DATE: 2003-01-31
; NUMBER OF SEQ ID NOS: 997516
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 133076
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapien

US-10-355-577-133076
Query Match          71.1%; Score 12.8; DB 14; Length 25;
Best Local Similarity 68.8%; Pred. No. 3.5e+03;
Matches 11; Conservative 3; Mismatches 2; Indels 0; Gaps 0;

QY          2 AUGGCCCUAUCGGUGCG 17
          | | | | : | : | : | : |
Db          5 AAGGACTATCGGTGCG 20

RESULT 15
US-10-355-577-152466/c
; Sequence 152466, Application US/10355577
; GENERAL INFORMATION:
; APPLICANT: Mittmann, Michael
; TITLE OF INVENTION: Methods of Genetic Analysis of Probes: HG-U133
; FILE REFERENCE: 3121
; CURRENT APPLICATION NUMBER: US/10/355,577
; CURRENT FILING DATE: 2003-01-31
; NUMBER OF SEQ ID NOS: 997516
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 152466
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapien
US-10-355-577-152466
Query Match          71.1%; Score 12.8; DB 14; Length 25;
Best Local Similarity 62.5%; Pred. No. 3.5e+03;
Matches 10; Conservative 4; Mismatches 2; Indels 0; Gaps 0;

QY          3 UGGCCCUAUCGGUGCGA 18
          : | | : | : | : | : |
Db          21 TTGACTATCGGTGCGA 6

Search completed: July 6, 2003, 16:49:14
Job time : 625.545 secs
```